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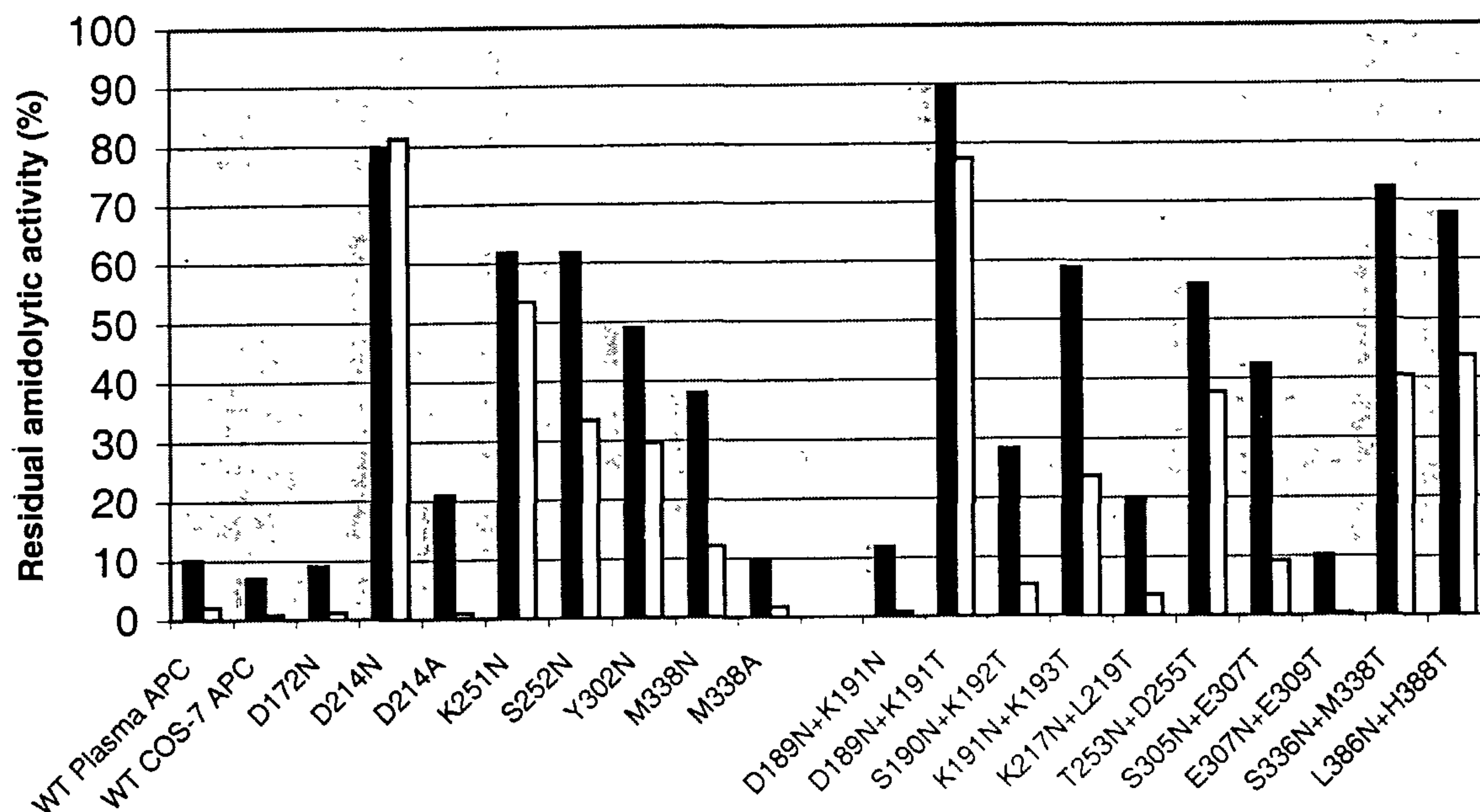
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(54) Title: PROTEIN C OR ACTIVATED PROTEIN C-LIKE MOLECULES



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The present invention relates to novel conjugates between polypeptide variants of pro-tein C and a non-polypeptide moiety, such as PEG or sugar moieties. In particular, the present invention provides novel protein C conjugates having an increased resistance to inactivation by e.g. human plasma and α 1-antitrypsin. Consequently, such conjugates have an increased in vivo half-life. Preferred examples include protein C conjugates, wherein at least one additional in vivo N-glycosylation site has been introduced. The conjugates of the invention are useful for treating a variety of diseases, including septic shock.

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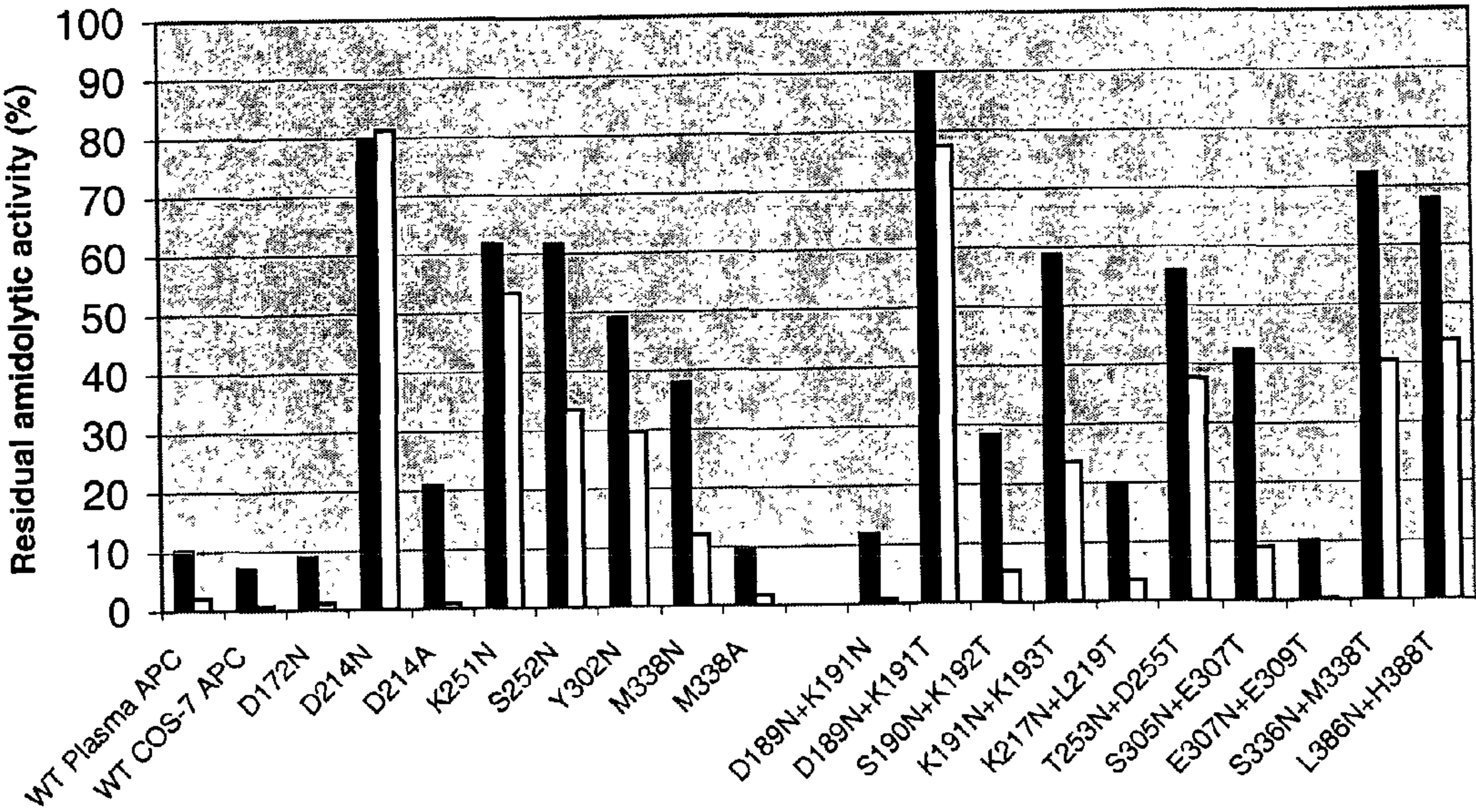
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(54) Title: PROTEIN C OR ACTIVATED PROTEIN C-LIKE MOLECULES



(57) Abstract: The present invention relates to novel conjugates between polypeptide variants of pro-teín C and a non-polypeptide moiety, such as PEG or sugar moieties. In particular, the present invention provides novel protein C conjugates having an increased resistance to inactivation by e.g. human plasma and α 1-antitrypsin. Consequently, such conjugates have an increased in vivo half-life. Preferred examples include protein C conjugates, wherein at least one additional in vivo N-glycosylation site has been introduced. The conjugates of the invention are useful for treating a variety of diseases, including septic shock.

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PROTEIN C OR ACTIVATED PROTEIN C-LIKE MOLECULES

FIELD OF THE INVENTION

The present invention relates to novel conjugates between polypeptide variants of protein C and a non-polypeptide moiety, to means and methods for preparing such conjugates, to pharmaceutical compositions comprising such conjugates and the use of such conjugates in therapy, in particular for the treatment of a variety of coagulation disorders. The present invention also relates to the polypeptide part of the conjugates of the invention.

BACKGROUND OF THE INVENTION

Blood coagulation is a process consisting of a complex interaction of various blood components, or factors, which eventually give rise to a fibrin clot. Generally, blood components participating in the coagulation "cascade" are proenzymes or zymogens, i.e. enzymatically inactive proteins that are converted into an active form by action of an activator. Regulation of blood coagulation is largely accomplished enzymatically by proteolytic inactivation of the pro-coagulation factors Va and VIIIa achieved by activated protein C (APC) (Esmon, J Biol Chem 1989; 264; 4743-4746).

Protein C is a serine protease that circulates in the plasma as a zymogen with a half-life of approximately 7 hours and plasma levels are typically in the range of 3-5 mg/l. It is produced *in vivo* in the liver as a single chain precursor polypeptide of 461 amino acids. This polypeptide undergoes multiple post-translational modifications including a) cleavage of a 42 amino acid signal sequence; b) cleavage of lysine and arginine residues (positions 156 and 157) to make a two-chain inactive zymogen (a 155 amino acid light chain attached via a disulfide bridge to a 262 amino acid heavy chain); c) vitamin K-dependent carboxylation of nine glutamic acid residues of the light chain resulting in nine gamma-carboxyglutamic acid residues in the N-terminal region of the light chain; and d) carbohydrate attachment at four sites (one in the light chain and three in the heavy chain). Finally, the two-chain zymogen may be activated by removal of a dodecapeptide (the activation peptide) at the N-terminus of the heavy chain (positions 158-169) producing the activated protein C (APC).

Protein C is activated by limited proteolysis by thrombin in complex with thrombomodulin on the luminal surface of the endothelial cell. As explained above, activation liberates a small 12 amino acid peptide (designated the activation peptide) from the N-terminal of the heavy chain. The APC has a half-life of approximately 15 minutes in plasma.

In the presence of its cofactor, protein S, APC proteolytically inactivates factors Va and VIIIa, thereby reducing thrombin generation (Esmon, *Thromb Haemost* 1993; 70; 29-35). Protein S circulates reversibly bound to another plasma protein, C4b-binding protein. Only free protein S serves as a cofactor for APC. Since C4b-binding protein is an acute phase reactant, the plasma levels of this protein varies greatly in many diseases and thus influence the anticoagulant activity of the protein C system.

The gene encoding human protein C maps to chromosome 2q13-q14 (Patraccchini et al., *Hum Genet* 1989; 81; 191-192) spans over 11 kb, and comprises a coding region (exons II to IX) and a 5' untranslatable region encompassing exon I. The protein domains encoded by exons II to IX show considerable homology with other vitamin K-dependent coagulation proteins such as factor IX and X. Exon II codes for a signal peptide, while exon III codes for a propeptide and a 38 amino acid sequence containing 9 Glu residues. The propeptide contains a binding site for the carboxylase transforming the Glu residues into dicarboxylic acid (Gla) able to bind calcium ions, a step required for phospholipid binding and protein C anticoagulant activity (Cheung et al., *Arch Biochem Biophys* 1989; 274; 574-581). Exons IV, V and VI encodes a short connection sequence and two EGF-like domains, respectively. Exon VII encodes both a domain encompassing a 12 amino acid activation peptide released after activation of protein C by thrombin, and the dipeptide 156-157 which, when cleaved off, yields the mature two-chain form of the protein. Exons VIII and IX encodes the serine protease domain.

The complete amino acid sequence of the human protein C has been reported by Foster et al., *PNAS. USA* 1986; 82; 4673-4677 and includes a signal peptide, a propeptide, a light chain, a heavy chain and an activation peptide.

Protein C binds to the endothelial cell protein receptor (EPCR). Binding of APC to EPCR renders APC incapable of inactivating factor Va and VIIIa, whereas binding of protein C to EPCR apparently enhances the activation rate of protein C by the thrombin-thrombomodulin complex. The physiological importance of these interactions is presently unknown. Apparently the binding of protein C to EPCR is strictly dependent on the presence of the Gla domain in a phospholipid independent manner (Esmon et al., *Haematologica* 1999; 84; 363-368).

APC is inhibited in the plasma by the protein C inhibitor as well as by alpha-1-antitrypsin and alpha-2-macroglobulin.

The experimental three-dimensional structure of human APC has been determined to 2.8 Å resolution and reported by Mather et al., *EMBO J* 1996; 15; 6822-6831. They report the

X-ray structure of APC in a Gla-domainless form. The structure includes a covalently bound inhibitor (D-Phe-Pro-Arg chloromethylketone, PPACK).

Protein C is currently isolated from prothrombin concentrates produced by monoclonal antibody affinity chromatography. Furthermore, protein C is produced recombinantly by expression from mammalian cells or modified protein C.

APC is used for the treatment of genetic and acquired protein C deficiency and is suggested to be used as anticoagulant in patients with some forms of Lupus, following stroke or myocardial infarction, after venous thrombosis, disseminated intravascular coagulation (DIC), septic shock, emboli such as pulmonary emboli, transplantation, such as bone marrow transplantation, burns, pregnancy, major surgery/trauma and adult respiratory stress syndrome (ARDS).

Recombinant APC is produced by Eli Lilly and Co and phase III trials for the treatment of sepsis (Bernard et al., N Engl J Med (2001), 344, pp. 699-709) has recently been completed. Patients suffering from severe sepsis were given doses of 24 µg/kg/h for a total duration of 96 hours as infusion.

However, relatively high doses and frequent administration is necessary to reach and sustain the desired therapeutic or prophylactic effects of APC due to its short half-life. As a consequence adequate dose regulation is difficult to obtain and the need of frequent intravenous administrations of high levels of APC is problematic and expensive.

A molecule with a longer circulation half-life would decrease the number of necessary administrations and potentially provide more optimal therapeutic APC levels with concomitant enhanced therapeutic effect.

The circulation half-life of APC may be increased, e.g. as a consequence of reduced renal clearance, of reduced proteolytic degradation or reduced inhibition. This may be achieved, e.g., by conjugation APC to a non-polypeptide moiety, e.g. PEG or carbohydrates, capable of conferring a reduced renal clearance to the protein and/or effectively blocking proteolytic enzymes or inhibitors from physical contact with the protein. Furthermore, this may also be achieved by mutating the protein C molecule in such a way that it remains active but blocks the binding of inhibitors to the protein.

PEGylated wild-type APC is described in JP 8-92294.

WO 91/09960 discloses a hybrid protein comprising modifications in the heavy chain part of protein C.

WO 01/59084 describes protein C variants comprising the substitutions D167F+D172K in combination with at least one further substitution in position 10, 11, 12, 32, 194, 195, 228, 149, 254, 302 or 316. The variants disclosed in WO 01/59084 are stated to have an increased anticoagulant activity.

5 WO 98/44000 broadly describes protein C variants with an increased amidolytic activity.

EP 0 323 149 describes zymogen forms of protein C with the following mutations in the heavy chain: D167F/G/Y/W. Such variants are stated to have an increased sensitivity to activation by thrombin.

10 WO 00/66754 reported that substitution of the residues naturally occurring in the positions 194, 195, 228, 249, 254, 302 or 316 lead to an increased half-life of APC in human blood as compared to the wild-type APC. The variants disclosed in WO 00/66754 are not within the scope of the present invention.

WO 99/63070 describes a C-terminally truncated form of protein C.

15 EP 0 946 715 reported chimeric protein C polypeptides where the protein C Gla domain was replaced by Gla domains from other vitamin K-dependent polypeptides, such as factor VII, factor X and prothrombin.

WO 99/20767 and WO 00/66753 discloses vitamin K-dependent polypeptide variants containing modifications in the Gla domain.

20 US 5,453,373 discloses human protein C derivatives which have altered glycosylation patterns and altered activation regions, such as N313Q and N329Q. The variants disclosed in US 5,453,373 are not within the scope of the present invention.

US 5,460,953 discloses DNA sequences encoding zymogen forms of protein C, which have been engineered so that one or more of the naturally occurring glycosylation sites have
25 been removed. More specifically, US 5,460,953 discloses the variants N97Q, N248Q, N313Q and N329Q. The variants disclosed in US 5,460,953 are not within the scope of the present invention. None of the disclosed variants in any of the above-mentioned prior art references are within the scope of the present invention.

US 5,270,178 is directed to specific protein C variants, wherein I 171 is deleted and
30 wherein Asp is replaced by Asn.

US 5,041,376 relates to a method for identifying and shielding functional sites or epitopes of transportable proteins, wherein additional N-linked glycosylation site(s) have been introduced.

US 5,766,921 is directed to protein C variants having increased resistance to inactivation by human plasma or α_1 -antitrypsin, where the heavy chain contains substitutions from the corresponding bovine heavy chain.

WO 01/57193 reports a protein C variant comprising a double mutation, one mutation
5 in positions 10, 11, 32 or 33 and one mutation in positions 194, 195, 228, 249, 254, 392 or 316.

WO 01/36462 relates to protein C variants comprising a substitution in position 12, optionally combined with substitutions in positions 10 and/or 11.

WO 00/26354 is directed to a method for producing glycosylated protein variants having reduced allergenicity.

10 WO 00/26230 is directed to a method for selecting a protein variant having reduced immunogenicity.

The DNA sequence and the corresponding amino acid sequence of human wild-type protein C, including the precursor form thereof, is disclosed in *inter alia* US 4,775,624 and US 4,968,626.

15 None of the variants disclosed in any of the above-identified patents/patent applications are within the scope of the present invention.

BRIEF DISCLOSURE OF THE INVENTION

The present invention relates to novel conjugates between polypeptide variants of protein C and a non-polypeptide moiety, to means and methods for preparing such conjugates, to
20 pharmaceutical compositions comprising such conjugates and the use of such conjugates in therapy, in particular for the treatment of a variety of coagulation disorders. The present invention also relates to the polypeptide part of the conjugates of the invention.

Accordingly, in its first aspect the invention relates to a conjugate comprising at least
25 one non-polypeptide moiety covalently attached to a protein C polypeptide that comprises an amino acid sequence which differs from that of a parent protein C polypeptide in at least one introduced and/or at least one removed amino acid residue comprising an attachment group for said non-polypeptide moiety.

In a further aspect the invention relates to a variant of a parent protein C polypeptide,
30 said variant comprising a substitution in a position selected from the group consisting of D172, D189, S190, K191, K192, K193, D214, E215, S216, K217, K218, L220, V243, V245, S250, K251, S252, T253, T254, D255, L296, Y302, H303, S304, S305, R306, E307, K308, E309, A310, R312, T315, F316, V334, S336, N337, M338, I348, L349, D351, R352, E357, E382,

G383, L386, L387 and H388, with the proviso that the substitution is not selected from the group consisting of T254S, T254A, T254H, T254K, T254R, T254N, T254D, T254E, T254G, T254Q, Y302S, Y302A, Y302T, Y302H, Y302K, Y302R, Y302N, Y302D, Y302E, Y302G, Y302Q, F316S, F316A, F316T, F316H, F316K, F316R, F316N, F316D, F316E, F316G and
5 F316Q.

In an even further aspect, the present invention relates to the polypeptide part of the conjugate of the invention.

In still further aspects the present invention relates to a nucleotide sequence encoding the polypeptide part of the conjugate of the invention, to a nucleotide sequence encoding the
10 polypeptide variant of the invention, to an expression vector comprising the nucleotide sequence of the invention and to a host cell comprising the nucleotide sequence of the invention or comprising the expression vector of the invention

Still other aspects of the present invention relates to a pharmaceutical composition comprising the conjugate or the variant of the invention as well as to methods of producing and
15 using the conjugates and variants of the invention.

DETAILED DISCLOSURE OF THE INVENTION

Definitions

In the context of the present application and invention the following definitions apply:

20 The term "conjugate" (or interchangeably "conjugated polypeptide") is intended to indicate a heterogenous (in the sense of composite or chimeric) molecule formed by the covalent attachment of one or more polypeptides to one or more non-polypeptide moieties such as polymer molecules, lipophilic compounds, sugar moieties or organic derivatizing agents. Preferably, the conjugate is soluble at relevant concentrations and conditions, i.e. soluble in
25 physiological fluids such as blood. Examples of conjugated polypeptides of the invention include glycosylated polypeptides and PEGylated polypeptides.

The term "covalent attachment" or "covalently attached" means that the polypeptide and the non-polypeptide moiety are either directly covalently joined to one another or are indirectly covalently joined to one another through an intervening moiety or moieties such as a
30 bridge, spacer or linkage moiety or moieties.

The term "non-conjugated polypeptide" may be used about the polypeptide part of the conjugate.

The term "non-polypeptide moiety" is intended to mean a molecule, different from a peptide polymer composed of amino acid monomers and linked together by peptide bonds, which molecule is capable of conjugating to an attachment group of the polypeptide of the invention. Preferred examples of such molecules include polymer molecules, sugar moieties, lipophilic compounds or organic derivatizing agents. When used in the context of a conjugate of the invention it will be understood that the non-polypeptide moiety is linked to the polypeptide part of the conjugate through an attachment group of the polypeptide. As explained above, the non-polypeptide moiety can be directly covalently joined to the attachment group or it can be indirectly covalently joined to the attachment group through an intervening moiety or moieties, such as a bridge spacer or linker moiety or moieties.

The term "polymer molecule" is a molecule formed by covalent linkage of two or more monomers, wherein none of the monomers is an amino acid residue, except where the polymer is human albumin or another abundant plasma protein. The term "polymer" can be used interchangeably with the term "polymer molecule" or "polymeric group".

The term "sugar moiety" is intended to indicate a carbohydrate-containing molecule comprising one or more monosaccharide residues, capable of being attached to the polypeptide (to produce a polypeptide conjugate in the form of a glycosylated polypeptide) by way of *in vivo* or *in vitro* glycosylation. The term "*in vivo* glycosylation" is intended to mean any attachment of a sugar moiety occurring *in vivo*, i.e. during posttranslational processing in a glycosylating cell used for expression of the polypeptide, e.g. by way of N-linked and O-linked glycosylation. The exact oligosaccharide structure depends, to a large extent, on the glycosylating organism in question. The term "*in vitro* glycosylation" is intended to refer to a synthetic glycosylation produced *in vitro*, normally involving covalently linking a sugar moiety to an attachment group of a polypeptide, optionally using a cross-linking agent. *In vivo* and *in vitro* glycosylation are discussed in detail further below.

An "N-glycosylation site" has the sequence N-X-S/T/C", wherein X is any amino acid residue except proline, N is asparagine and S/T/C is either serine, threonine or cysteine, preferably serine or threonine, and most preferably threonine. An "O-glycosylation site" is the OH-group of a serine or threonine residue.

The term "attachment group" is intended to indicate a functional group of the polypeptide, in particular of an amino acid residue thereof or a carbohydrate moiety, capable of attaching a non-polypeptide moiety such as a polymer molecule, a sugar moiety, a lipophilic molecule

or an organic derivatizing agent. Useful attachment groups and their matching non-polypeptide moieties are apparent from the table below.

Attachment group	Amino acid	Examples of non-polypeptide moiety	Conjugation method/- Activated PEG	Reference
-NH ₂	N-terminal, Lys	Polymer, e.g. PEG, with amide or imine group	mPEG-SPA Tresylated mPEG	Shearwater Inc. Delgado et al., critical reviews in Therapeutic Drug Carrier Systems 9(3,4):249-304 (1992)
-COOH	C-terminal, Asp, Glu	Polymer, e.g. PEG, with ester or amide group Oligosaccharide moiety	mPEG-Hz <i>In vitro</i> coupling	Shearwater Inc.
-SH	Cys	Polymer, e.g. PEG, with disulfide, maleimide or vinyl sulfone group Oligosaccharide moiety	PEG- vinylsulphone PEG-maleimide <i>In vitro</i> coupling	Shearwater Inc. Delgado et al., critical reviews in Therapeutic Drug Carrier Systems 9(3,4):249-304 (1992)
-OH	Ser, Thr, OH-, Lys	Oligosaccharide moiety PEG with ester, ether, carbamate, carbonate	<i>In vivo</i> O-linked glycosylation	

-CONH ₂	Asn as part of an N-glycosylation site	Oligosaccharide moiety Polymer, e.g. PEG	<i>In vivo</i> N-glycosylation	
Aromatic residue	Phe, Tyr, Trp	Oligosaccharide moiety	<i>In vitro</i> coupling	
-CONH ₂	Gln	Oligosaccharide moiety	<i>In vitro</i> coupling	Yan and Wold, Biochemistry, 1984, Jul 31; 23(16): 3759-65
Aldehyde Ketone	Oxidized oligo-saccharide	Polymer, e.g. PEG, PEG-hydrazide	PEGylation	Andresz et al., 1978, Makromol. Chem. 179:301, WO 92/16555, WO 00/23114
Guanidino	Arg	Oligosaccharide moiety	<i>In vitro</i> coupling	Lundblad and Noyes, Chemical Reagents for Protein Modification, CRC Press Inc., Florida, USA
Imidazole ring	His	Oligosaccharide moiety	<i>In vitro</i> coupling	As for guanidine

For *in vivo* N-glycosylation, the term “attachment group” is used in an unconventional way to indicate the amino acid residues constituting an N-glycosylation site. Although the asparagine residue of the N-glycosylation site is the one to which the sugar moiety is attached during glycosylation, such attachment cannot be achieved unless the other amino acid residues of the N-glycosylation site are present.

Accordingly, when the non-polypeptide moiety is a sugar moiety and the conjugation is to be achieved by N-glycosylation, the term “amino acid residue comprising an attachment group for the non-polypeptide moiety” as used in connection with alterations of the amino acid sequence of the polypeptide of interest is to be understood as meaning that one or more amino

acid residues constituting an N-glycosylation site are to be altered in such a manner that either a functional N-glycosylation site is introduced into the amino acid sequence or removed from said sequence.

Amino acid names and atom names (e.g. CA, CB, CD, CG, SG, NZ, N, O, C, etc.) are used as defined by the Protein DataBank (PDB) (www.pdb.org), which is based on the IUPAC nomenclature (IUPAC Nomenclature and Symbolism for Amino Acids and Peptides (residue names, atom names, etc.), *Eur. J. Biochem.*, 138, 9-37 (1984) together with their corrections in *Eur. J. Biochem.*, 152, 1 (1985)).

The term "amino acid residue" is intended to indicate an amino acid residue contained in the group consisting of alanine (Ala or A), cysteine (Cys or C), aspartic acid (Asp or D), glutamic acid (Glu or E), phenylalanine (Phe or F), glycine (Gly or G), histidine (His or H), isoleucine (Ile or I), lysine (Lys or K), leucine (Leu or L), methionine (Met or M), asparagine (Asn or N), proline (Pro or P), glutamine (Gln or Q), arginine (Arg or R), serine (Ser or S), threonine (Thr or T), valine (Val or V), tryptophan (Trp or W) and tyrosine (Tyr or Y) residues.

The terminology used for identifying amino acid positions/substitutions is illustrated as follows: K174 in a given amino acid sequence indicates that position number 174 is occupied by a lysine residue in the amino acid sequence shown in SEQ ID NO:2 or 4. K174S indicates that the lysine residue of position 174 is substituted with a serine residue. Alternative substitutions are indicated with a "/", e.g., K174S/T means that the lysine residue of position 174 is substituted with either a serine residue or a threonine residue. Multiple substitutions are indicated with a "+", e.g., D172N+K174S means that the aspartic acid residue of position 172 is substituted with an asparagine residue and that the lysine residue in position 174 is substituted with a serine residue. The insertion of an additional amino acid residue is indicated in the following way: Insertion of an alanine residue after K174 is indicated by K174KA. A deletion of an amino acid residue is indicated by an asterisk. For example, deletion of the lysine residue of position 174 is indicated by K174*. Unless otherwise indicated, the numbering of amino acid residues made herein is made relative to the amino acid sequence of SEQ ID NO:2 or 4.

The term "differs" or "differs from" when used in connection with specific mutations is intended to allow for additional differences being present apart from the specified amino acid difference. For instance, in addition to the removal and/or introduction of amino acid residues comprising an attachment group for the non-polypeptide moiety the protein C polypeptide can comprise other substitutions, insertions or deletions, which are not related to the introduction/removal of such amino acid residues. Thus, in addition to the amino acid alterations dis-

closed herein aimed at removing and/or introducing attachment sites for the non-polypeptide moiety, it will be understood that the amino acid sequence of the polypeptide conjugate of the invention may, if desired, contain other alterations that need not be related to introduction or removal of attachment sites, i.e. other substitutions, insertions or deletions. These may, for example, include truncation of the N- and/or C-terminus by one or more amino acid residues, or addition of one or more extra residues at the N- and/or C-terminus, e.g. addition of a methionine residue at the N-terminus as well as “conservative amino acid substitutions”, i.e. substitutions performed within groups of amino acids with similar characteristics, e.g. small amino acids, acidic amino acids, polar amino acids, basic amino acids, hydrophobic amino acids and aromatic amino acids.

Examples of conservative substitutions in the present invention may in particular be selected from the groups listed in the table below.

1	Alanine (A)	Glycine (G)	Serine (S)	Threonine (T)
2	Aspartic acid (D)	Glutamic acid (E)		
3	Asparagine (N)	Glutamine (Q)		
4	Arginine (R)	Histidine (H)	Lysine (K)	
5	Isoleucine (I)	Leucine (L)	Methionine (M)	Valine (V)
6	Phenylalanine (F)	Tyrosine (Y)	Tryptophan (W)	

When used in the present context the term “precursor protein C” refers to the DNA-encoded form of protein C, i.e. it includes the signal peptide (residue -42 to -1), the light chain (residue 1-155), the Lys-Arg dipeptide (residue 156-157) and the heavy chain (158-419), including the activation peptide (residue 158-169), shown in SEQ ID NO:2.

The term “two-chain zymogen protein C” refers to the secreted, inactive form of protein C, which includes the light chain (residue 1-155) and the heavy chain (158-419), including the activation peptide (158-169), shown in SEQ ID NO:4.

The term “one-chain zymogen protein C” refers to the inactive form of protein C, which includes the light chain (residue 1-155), the heavy chain (158-419), including the activation peptide (158-169), and the Lys-Arg dipeptide (residue 156-157) shown in SEQ ID NO:4.

Whenever the term “zymogen protein C” is used this term refers to both the one-chain form and the two-chain form of the zymogen protein C.

The terms “activated protein C”, “activated human protein C”, “APC” or “human APC” are used about the activated zymogen and includes the light chain (residue 1-155) and the heavy chain (without the activation peptide) of SEQ ID NO:4. The latter amino acid sequence, i.e. the amino acid sequence of activated protein C is sometimes referred to herein as “the APC
5 part of the amino acid sequence shown in SEQ ID NO:4”.

The term “protein C” encompasses all of the above-mentioned forms of protein C, i.e. the “precursor protein C” form, the “zymogen protein C” form (the one-chain form as well as the two-chain form) and the “activated protein C form”.

The term “parent” is intended to indicate the molecule to be improved in accordance
10 with the present invention. Although the parent polypeptide to be modified by the present invention may be any protein C polypeptide, and thus be derived from any origin, e.g. a non-human mammalian origin, it is preferred that the parent polypeptide is human protein C (i.e. human precursor protein C, human zymogen protein C or human activated protein C) or a fragment or variant thereof.

15 A fragment is a part of the full-length human protein C sequence, e.g. a C-terminally or N-terminally truncated version thereof. Specific examples of parent protein C polypeptide fragments include human protein C terminally truncated with 1-15 amino acid residues and/or N-terminally truncated with 1-3 amino acid residues.

As mentioned above, the parent protein C polypeptide may also be a variant of human
20 protein C. Specific examples of variants of human protein C includes e.g. addition of a methionine residue at the N-terminus as well as variants containing one or more conservative amino acid substitutions as discussed above. Other examples of variants include human protein C variants wherein one or more amino acids in the protein C Gla domain has been substituted or wherein the entire protein C Gla domain has been substituted with another Gla domain, e.g.
25 the Gla domain of protein S.

The term “variant” (of a parent polypeptide) is intended to cover a polypeptide, which differs in one or more amino acid residues from its parent polypeptide, normally in 1-15 amino acid residues (such as in 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15 amino acid residues), e.g. in 1-10 amino acid residues or in 1-5 amino acid residues.

30 The term “mutation” and “substitution” are used interchangeably herein.

The term “nucleotide sequence” is intended to indicate a consecutive stretch of two or more nucleotide molecules. The nucleotide sequence may be of genomic, cDNA, RNA, semi-synthetic or synthetic origin, or any combination thereof.

The term “polymerase chain reaction” or “PCR” generally refers to a method for amplification of a desired nucleotide sequence *in vitro* as described, for example, in US 4,683,195. In general, the PCR method involves repeated cycles of primer extension synthesis, using oligonucleotide primers capable of hybridising preferentially to a template nucleic acid.

5 “Cell”, “host cell”, “cell line” and “cell culture” are used interchangeably herein and all such terms should be understood to include progeny resulting from growth or culturing of a cell.

“Transformation” and “transfection” are used interchangeably to refer to the process of introducing DNA into a cell.

10 “Operably linked” refers to the covalent joining of two or more nucleotide sequences, by means of enzymatic ligation or otherwise, in a configuration relative to one another such that the normal function of the sequences can be performed. For example, the nucleotide sequence encoding a presequence or secretory leader is operably linked to a nucleotide sequence coding for a polypeptide if it is expressed as a preprotein that participates in the secretion of the polypeptide: a promoter or enhancer is operably linked to a coding sequence if it affects the tran-
15 scription of the sequence; a ribosome binding site is operably linked to a coding sequence if it is positioned so as to facilitate translation. Generally, “operably linked” means that the nucleotide sequences being linked are contiguous and, in the case of a secretory leader, contiguous and in reading phase. Linking is accomplished by ligation at convenient restriction sites. If such sites
20 do not exist, then synthetic oligonucleotide adaptors or linkers are used, in conjunction with standard recombinant DNA methods.

The term “introduce” is primarily intended to mean substitution of an existing amino acid residue, but may also mean insertion of an additional amino acid residue.

25 The term “remove” is primarily intended to mean substitution of the amino acid residue to be removed by another amino acid residue, but may also mean deletion (without substitution) of the amino acid residue to be removed.

The term “functional *in vivo* half-life” is used in its normal meaning, i.e. the time at which 50% of the biological activity of the polypeptide or conjugate is still present in the body/target organ, or the time at which the activity of the polypeptide or conjugate is 50% of
30 the initial value. As an alternative to determining functional *in vivo* half-life, “serum half-life” may be determined, i.e. the time at which 50% of the polypeptide or conjugate molecules circulate in the plasma or bloodstream prior to being cleared. Determination of serum half-life is often more simple than determining the functional *in vivo* half-life and the magnitude of serum

half-life is usually a good indication of the magnitude of functional *in vivo* half-life. Alternatively terms to serum half-life include “plasma half-life”, “circulating half-life”, “serum clearance”, “plasma clearance” and “clearance half-life”. The polypeptide or conjugate is cleared by the action of one or more of the reticuloendothelial systems (RES), kidney, spleen or liver, by tissue factor, SEC receptor or other receptor mediated elimination, or by specific or unspecific proteolysis. Normally, clearance depends on size (relative to the cutoff for glomerular filtration), charge, attached carbohydrate chains, and the presence of cellular receptors for the protein. The functionality to be retained is normally selected from anticoagulant, amidolytic or receptor binding activity. The functional *in vivo* half-life and the serum half-life may be determined by any suitable method known in the art.

The term “increased” as used about the functional *in vivo* half-life or serum half-life is used to indicate that the relevant half-life of the conjugate or polypeptide is statistically significantly increased relative to that of a reference molecule, e.g. APC, determined under comparable conditions. Normally, functional *in vivo* or serum half-life is increased when clearance, proteolytic degradation and/or inhibition of the polypeptide is decreased. Thus, preferred conjugates are such conjugates, which, in their activated form, has an increased functional *in vivo* half-life or an increased serum half-life as compared to human APC. Particular preferred conjugates are such conjugates where the ratio between the serum half-life (or functional *in vivo* half-life) of said conjugate and the serum half-life (or functional *in vivo* half-life) of human APC is at least 1.25, more preferably at least 1.50, such as at least 1.75, e.g. at least 2, even more preferably at least 3, such as at least 4, e.g. at least 5, most preferably at least 6, such as at least 7, e.g. at least 8, at least 9 or at least 10.

Clearance mechanisms of relevance for a polypeptide or conjugate of the invention may include one or more of the reticuloendothelial systems (RES), kidney, spleen or liver, receptor-mediated degradation, or specific or non-specific proteolysis. The term “renal clearance” is used in its normal meaning to indicate any clearance taking place by the kidneys, e.g. by glomerular filtration, tubular excretion or tubular elimination. Normally, renal clearance depends on physical characteristics of the conjugate, including molecular weight, size (relative to the cutoff for glomerular filtration), symmetry, shape/rigidity and charge. A molecular weight of about 67 kDa is normally considered to be a cut-off-value for renal clearance. Renal clearance may be measured by any suitable assay, e.g. an established *in vivo* assay. For instance, renal clearance may be determined by administering a labelled (e.g. radiolabelled or fluorescence labelled) polypeptide conjugate to a patient and measuring the label activity in urine col-

lected from the patient. Reduced renal clearance is determined relative to the reference molecule, such as APC.

The term “activity”, “APC activity” or “activated protein C activity” is intended to indicate that the conjugate of the invention, in its activated form, retain the essential properties of APC.

A suitable *in vitro* APC activity assay (entitled “APC Amidolytic Assay”) is described in Example 9 herein. Thus, more particularly, a conjugate of the present invention is classified as having “APC activity” if the conjugate, in its activated form, has an activity of at least 10% of the human APC activity when tested in the “APC Amidolytic Assay” described in Example 9 herein. Preferably, the conjugate has an activity of at least 20% of the human APC activity, such as an activity of at least 30% of the human APC activity, more preferably the conjugate has an activity of at least 40% of the human APC activity, such as an activity of at least 50% of the human APC activity, even more preferably the conjugate has an activity of at least 60% of the human APC activity, e.g. an activity of at least 70% of the human APC activity, most preferably the conjugate has an activity of at least 80% of the human APC activity, such as an activity of at least 90% of the human APC activity. In a very interesting embodiment, the conjugate has an activity, when tested in the “APC Amidolytic Assay” described in Example 9 herein, which is essentially the same or higher than the activity of human APC. It will be understood that the conjugate of the invention and the wild-type human APC should be assayed under identical conditions, i.e. the concentration of both proteins should be identical when assayed as described in Example 9 herein.

Alternatively, the “APC activity” may be measured in the *in vitro* assay entitled “APC Clotting Assay” described in Example 10 herein. More particularly, a conjugate of the present invention is classified as having “APC activity” if the conjugate, in its activated form, has an anticoagulant activity of at least 5% of the human APC anticoagulant activity when tested in the “APC Clotting Assay” described in Example 10 herein. Preferably, the conjugate has an anticoagulant activity of at least 10% of the human APC anticoagulant activity, such as an anticoagulant activity of at least 20% of the human APC anticoagulant activity, e.g. an anticoagulant activity of at least 30%, more preferably the conjugate has an anticoagulant activity of at least 40% of the human APC anticoagulant activity, such as an anticoagulant activity of at least 50% of the human APC anticoagulant activity, even more preferably the conjugate has an anticoagulant activity of at least 60% of the human APC anticoagulant activity, e.g. an anticoagulant activity of at least 70% of the human APC anticoagulant activity, most preferably the conjugate

has an anticoagulant activity of at least 80% of the human APC anticoagulant activity, such as an anticoagulant activity of at least 90% of the human APC anticoagulant activity. In a very interesting embodiment, the conjugate has an anticoagulant activity, when tested in the "APC Clotting Assay" described in Example 10 herein, which is essentially the same or higher than the anticoagulant activity of human APC. Examples of typical PC activity intervals are, for example 5-75% of the human APC anticoagulant activity, such as 10-50% of the human APC anticoagulant activity, such as 10-40% of the human APC anticoagulant activity. It will be understood that the conjugate of the invention and the wild-type human APC should be assayed under identical conditions, i.e. the concentration of both proteins should be identical when assayed as described in Example 10 herein.

The terms "increased resistance towards inactivation by alpha-1-antitrypsin" and "increased resistance towards inactivation by human plasma", respectively, are primarily intended to mean a conjugate of the invention which is inhibited by alpha-1-antitrypsin or human plasma, respectively, to a lesser degree than human APC. In order to enable the skilled person, at an early stage of his development work, to select effective and preferred conjugates, the present inventors have developed suitable preliminary tests, which can easily be carried out by the skilled person in order to initially assess the performance of the conjugate in question. Thus, the "Alpha-1-Antitrypsin Inactivation Assay" (described in Example 11 herein), the "Human Plasma Inactivation Assay I" (described in Example 12 herein) and the "Human Plasma Inactivation Assay II" (described in Example 13 herein) may be employed to initially assess the potential of a selected conjugate. Using either the first, the second, the third or all of these tests, the suitability of a selected conjugate to resist inactivation by either alpha-1-antitrypsin and/or human plasma can be assessed, the rationale being that if a conjugate is strongly inhibited by either alpha-1-antitrypsin or human plasma, or both, it is normally not necessary to carry out further test experiments.

Therefore, a conjugate, which is particularly interesting for the purposes described herein, is a conjugate which, in its activated form, has a residual activity of at least 20% when tested in the "Alpha-1-Antitrypsin Inactivation Assay" described in Example 11 herein using an inhibitor concentration of 16.6 μ M. Preferably, the conjugate has a residual activity of at least 30%, such as a residual activity of at least 40%, more preferably the conjugate has a residual activity of at least 50%, such as a residual activity of at least 60%, even more preferably the conjugate has a residual activity of at least 70%, such as a residual activity at least 75%, most preferably the conjugate has a residual activity of at least 80%, such as at least 85%.

Alternatively, or in addition to the above-mentioned test, the suitability of a selected conjugate may be tested in the "Human Plasma Inactivation Assay I". Thus, a conjugate which is particular interesting for the purposes described herein, is a conjugate which, in its activated form, has a residual activity of at least 20% when tested in the "Human Plasma Inactivation Assay I" described in Example 12 herein. Preferably, the conjugate has a residual activity of at least 30%, such as a residual activity of at least 40%, more preferably the conjugate has a residual activity of at least 50%, such as a residual activity of at least 60%, even more preferably the conjugate has a residual activity of at least 70%, such as a residual activity at least 75%.

Alternatively, or in addition to the above-mentioned test(s), the suitability of a selected conjugate may be tested in the "Human Plasma Inactivation Assay II". Thus, a conjugate which is particular interesting for the purposes described herein, is a conjugate where the ratio between the *in vitro* half-life of said conjugate, in its activated form, and the *in vitro* half-life of human APC is at least 1.25 when tested in the "Human Plasma Inactivation Assay II" described in Example 13 herein, preferably at least 1.5, such as at least 2, more preferably at least 3, such as at least 4, even more preferably at least 5, such as at least 6, most preferably at least 7, such as at least 8, in particular at least 9, such as at least 10.

The term "reduced immunogenicity" is intended to indicate that the conjugate gives rise to a measurably lower immune response than a reference molecule, e.g. wild-type human APC or wild-type human protein C, as determined under comparable conditions. The immune response may be a cell or antibody mediated response (see, e.g., Roitt: Essential Immunology (8th Edition, Blackwell) for further definition of immunogenicity). Normally, reduced antibody reactivity is an indication of reduced immunogenicity. Reduced immunogenicity may be determined by use of any suitable method known in the art, e.g. *in vivo* or *in vitro*.

The terms "at least 25% of its side chain exposed to the surface" and "at least 50% of its side chain exposed to the surface" are defined with reference to Example 1, where the calculations, etc. are described in detail.

Conjugate of the invention

The conjugates of the present invention are the result of a generally new strategy for developing improved protein C molecules. More specifically, by removing and/or introducing an amino acid residue comprising an attachment group for the non-polypeptide moiety it is possible to specifically adapt the polypeptide so as to make the molecule more susceptible to conjugation to the non-polypeptide moiety of choice, to optimize the conjugation pattern, e.g. to

ensure an optimal distribution and number of non-polypeptide moieties on the surface of the protein C molecule and to ensure that only the attachment groups intended to be conjugated is present in the molecule, and thereby obtain a new conjugate molecule, which has APC activity and in addition one or more improved properties as compared to protein C molecules available
5 today. For instance, when the total number of amino acid residues comprising an attachment group for the non-polypeptide of choice is increased or decreased to an optimized level, the renal clearance of the conjugate is typically significantly reduced due to the altered shape, size and/or charge of the molecule achieved by the conjugation. Furthermore, we have found that it is possible to design the attachment of a non-polypeptide moiety to an attachment group in the
10 polypeptide part of the conjugate so that inactivation by human plasma or certain inhibitors, such as alpha-1-antitrypsin, is significantly reduced (see below).

The amino acid residue comprising an attachment group for a non-polypeptide moiety, either it be removed or introduced, is selected on the basis of the nature of the non-polypeptide moiety of choice and, in most instances, on the basis of the method in which conjugation be-
15 tween the polypeptide and the non-polypeptide moiety is to be achieved. For instance, when the non-polypeptide moiety is a polymer molecule such as a polyethylene glycol or polyalkylene oxide derived molecule amino acid residues comprising an attachment group may be selected from the group consisting of lysine, cysteine, aspartic acid, glutamic acid, histidine, and tyro-
sine, preferably cysteine and lysine, in particular lysine. When the non-polypeptide moiety is a
20 sugar moiety the attachment group is, e.g., an *in vivo* glycosylation site, preferably an N-glycosylation site.

Whenever an attachment group for a non-polypeptide moiety is to be introduced into or removed from the protein C polypeptide in accordance with the present invention, the position of the polypeptide to be modified is conveniently selected as follows:

25 The position is preferably located at the surface of the protein C polypeptide, and more preferably occupied by an amino acid residue having more than 25% of its side chain exposed to the surface, such as more than 50% of its side chain exposed to the surface. Such positions have been identified on the basis of an analysis of a 3D structure of the human wild-type APC molecule as described in the Methods section herein. Furthermore, homologous positions in
30 non-human APC polypeptides (including variants thereof) comprising an amino acid sequence being homologous to that of wild-type human protein C may easily be determined by suitable alignment of the respective sequences or 3D structures.

In order to determine an optimal distribution of attachment groups, the distance between amino acid residues located at the surface of the polypeptide is calculated on the basis of a 3D structure of the polypeptide. More specifically, the distance between the CB's of the amino acid residues comprising such attachment groups, or the distance between the functional group (NZ for lysine, CG for aspartic acid, CD for glutamic acid, SG for cysteine) of one and the CB of another amino acid residue comprising an attachment group are determined. In case of glycine, CA is used instead of CB. In the polypeptide part of a conjugate of the invention, any of said distances is preferably more than 8 Å, in particular more than 10 Å in order to avoid or reduce heterogeneous conjugation.

The total number of amino acid residues to be altered in accordance with the present invention, e.g. as described in the subsequent sections herein, (as compared to the parent protein C molecule) will typically not exceed 15. The exact number of amino acid residues and the type of amino acid residues to be introduced depends, *inter alia*, on the desired nature and degree of conjugation (e.g. the identity of the non-polypeptide moiety, how many non-polypeptide moieties it is desirable or possible to conjugate to the polypeptide, where in the polypeptide conjugation should be performed or avoided, etc.). Preferably, the polypeptide part of the conjugate of the invention or the polypeptide of the invention comprises an amino acid sequence which differs in 1-15 amino acid residues from the amino acid sequence shown in SEQ ID NO:4, such as in 1-8 or 2-8 amino acid residues, e.g. in 1-5 or 2-5 amino acid residues. Thus, normally the polypeptide part of the conjugate or the polypeptide of the invention comprises an amino acid sequence which differs from the amino acid sequence shown in SEQ ID NO:4 in 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15 amino acid residues.

Preferably, the conjugate of the invention has one or more improved properties as compared to wild-type human APC, including increased functional *in vivo* half-life, increased serum half-life, increased resistance to inhibitors, reduced renal clearance, reduced immunogenicity and/or increased bioavailability. It is contemplated that a conjugate of the present invention offers a number of advantages over the currently available APC products, including longer duration between injections, administration of less protein, and fewer side effects. Furthermore, a reduced anticoagulant activity might be beneficial for reducing the risk of bleeding while maintaining the anti-inflammatory effect of the APC conjugates. This might be especially important when the conjugate has an extended plasma half-life. These new properties should enhance the anti-inflammatory effect compared to the anticoagulant activity allowing a more effective and safe treatment.

Typically, the conjugate according to the invention has a molecular weight of at least about 67 kDa, preferably at least about 70 kDa, although a lower molecular weight may also give rise to a reduced renal clearance. Polymer molecules, such as PEG, or introduced glycosylation sites have been found to be particularly useful for adjusting the molecular weight of the conjugate.

The conjugate of the invention comprises a sufficient number or type of non-polypeptide moieties to improve one or more of the above mentioned desired properties of the protein C polypeptide. Normally a conjugate of the invention comprises 1-10 first non-polypeptide moieties, in particular 1-8 or 1-5 first non-polypeptide moieties.

The conjugate of the invention may further comprise at least one second non-polypeptide moiety which is different from said first non-polypeptide moiety. For instance, the conjugate of the invention may comprise 1-10 second non-polypeptide moieties, in particular 1-8 or 1-5 second non-polypeptide moieties. For instance, when the first non-polypeptide moiety is a sugar moiety, in particular an *in vivo* attached sugar moiety, a second non-polypeptide moiety of interest could be a polymer of the PEG type. The *in vivo* attached sugar moiety may be attached to a naturally occurring *in vivo* glycosylation site of the polypeptide, or an introduced site.

In a very interesting embodiment of the invention the non-polypeptide moiety is introduced in the active site region of protein C, the rationale being that introduction of a non-polypeptide moiety or moieties in this particular region of the protein C molecule will impair binding of inhibitors (such as alpha-1-antitrypsin) to APC while still retaining a substantial APC activity. This, in turn, has the consequence that such conjugates will exhibit a significantly prolonged half-life compared to wild-type human APC since elimination of the inhibitor/APC complex via hepatic receptors is avoided or at least reduced. Selection of amino acid residues, which are located in the active site region of protein C is described in detail in Example 2 herein.

When used herein the term "active site region" is defined with reference to Example 2 herein, where the actual amino acid residues which constitute the active site region are shown.

In a particular preferred embodiment of the invention, an attachment group for a non-polypeptide moiety is introduced in a position of the active site region which is occupied by an amino acid residue having at least 25% of its side chain exposed to the surface (see Example 3 herein), i.e. an attachment group for a non-polypeptide moiety is introduced in a position selected from the group consisting of D172, D189, S190, K191, K192, K193, D214, E215, S216,

K217, K218, L220, V243, V245, N248, S250, K251, S252, T253, T254, D255, L296, Y302, H303, S304, S305, R306, E307, K308, E309, A310, K311, R312, N313, R314, T315, F316, V334, S336, N337, M338, I348, L349, D351, R352, E357, E382, G383, L386, L387 and H388 (H211 and C384 being excluded). Preferably, the introduced attachment group is an attachment
 5 group for a sugar moiety, in particular an *in vivo* N-glycosylation site (see the section entitled *Conjugate of the invention where the non-polypeptide moiety is a sugar moiety*).

Conjugate of the invention where the non-polypeptide moiety is a sugar moiety

As explained above, in a preferred embodiment the present invention relates to a con-
 10 jugate comprising at least one introduced glycosylation site, in particular an *in vivo* N-glycosylation site, covalently attached to a protein C polypeptide that comprises an amino acid sequence which differs from a parent protein C polypeptide, in particular from the amino acid sequence shown in SEQ ID NO:4 or a variant thereof, in at least one introduced glycosylation site.

15 Preferably, the glycosylation site is introduced in a position, which is occupied by an amino acid residue having at least 25% of its side chain exposed to the surface, such as at least 50% of its side chain exposed to the surface. Such amino acid residues are identified in Example 1 herein. It should be understood that when the term "at least 25% (or at least 50%) of its side chain exposed to the surface" is used in connection with introduction of an *in vivo* N-
 20 glycosylation site this term refers to the surface accessibility of the amino acid side chain in the position where the sugar moiety is actually attached. In many cases it will be necessary to introduce a serine or a threonine residue in position +2 relative to the asparagine residue to which the sugar moiety is actually attached (unless, of course, this position is already occupied by a serine or a threonine residue) and these positions, where the serine or threonine residues are
 25 introduced, are allowed to be buried, i.e. to have less than 25% of their side chains exposed to the surface.

In order to impair the binding of inhibitors, such as alpha-1-antitrypsin, to APC the glycosylation site is preferably introduced in a position which is within the active site region (defined in Example 2 herein) and which is occupied by an amino acid residue having at least
 30 25% of its side chain exposed to the surface (defined in Example 3 herein), i.e. the introduced *in vivo* N-glycosylation site is preferably selected from the group consisting of D172N+K174S, D172N+K174T, D189N+K191S, D189N+K191T, S190N+K192S, S190N+K192T, K191N+K193S, K191N+K193T, K192N+L194S, K192N+L194T, K193N+A195S,

K193N+A195T, D214N, D214N+S216T, E215N+K217S, E215N+K217T, S216N+K218S,
 S216N+K218T, K217N+L219S, K217N+L219T, K218N+L220S, K218N+L220T,
 L220N+R222S, L220N+R222T, V243N+V245S, V243N+V245T, V245N+P247S,
 V245N+P247T, S250N, S250N+S252T, K251N, K251N+T253S, S252N, S252N+T254S,
 5 T253N+D255S, T253N+D255T, T254N+N256S, T254N+N256T, D255N+D257S,
 D255N+D257T, L296N, L296N+T298S, Y302N, Y302N+S304T, H303N, H303N+S305T,
 S304N+R306S, S304N+R306T, S305N+E307S, S305N+E307T, R306N+K308S,
 R306N+K308T, E307N+E309S, E307N+E309T, K308N+A310S, K308N+A310T,
 E309N+K311S, E309N+K311T, A310N+R312S, A310N+R312T, R312N+R314S,
 10 R312N+R314T, T315N+V317S, T315N+V317T, F316N+L318S, F316N+L318T, V334N,
 V334N+S336T, S336N+M338S, S336N+M338T, V339S, V339T, M338N, M338N+S340T,
 I348N+G350S, I348N+G350T, L349N+D351S, L349N+D351T, D351N+Q353S,
 D351N+Q353T, R352N+D354S, R352N+D354T, E357N+D359S, E357N+D359T,
 G383N+G385S, G383N+G385T, L386N+H388S, L386N+H388T, L387N+N389S,
 15 L387N+N389T, H388N+Y390S and H388N+Y390T.

More preferably, the introduced *in vivo* N-glycosylation site is selected from the group consisting of S190N+K192S, S190N+K192T, K191N+K193S, K191N+K193T,
 D189N+K191S, D189N+K191T, D214N, D214N+S216T, K217N+L219S, K217N+L219T,
 K251N, K251N+T253S, S252N, S252N+T254S, T253N+D255S, T253N+D255T, Y302N,
 20 Y302N+S304T, S305N+E307S, S305N+E307T, E307N+E309S, E307N+E309T,
 S336N+M338S, S336N+M338T, V339S, V339T, M338N, M338N+S340T, G383N+G385S,
 G383N+G385T, L386N+H388S and L386N+H388T.

Even more preferably the introduced *in vivo* N-glycosylation site is selected from the group consisting of D189N+K191T, K191N+K193T, D214N, K251N, S252N, T253N+D255T,
 25 Y302N, S305N+E307T, S336N+M338T, V339T, M338N, G383N+G385T, and most preferably the introduced *in vivo* N-glycosylation site is selected from the group consisting of
 D189N+K191T, K191N+K193T, D214N, T253N+D255T, S305N+E307T, S336N+M338T,
 M338N, G383N+G385T and L386N+H388T. In a particular preferred embodiment the introduced *in vivo* N-glycosylation site is selected from the group consisting of D189N+K191T,
 30 D214N and L386N+H388T.

As explained above, the increased resistance towards inactivation by alpha-1-antitrypsin and/or human plasma may be determined and assessed by the "Alpha-1-Antritrypsin

Inactivation Assay”, the “Human Plasma Inactivation Assay I” or the “Human Plasma Inactivation Assay II” disclosed herein.

The conjugate of the invention may contain a single *in vivo* glycosylation site (in addition to the already present glycosylation sites at positions 97, 248, 313 and 329). However, in order to obtain efficient shielding of protease cleavage sites on the surface of the parent polypeptide and/or to efficiently impair inhibitor binding, it is often desirable that the polypeptide part of the conjugate comprises more than one *in vivo* glycosylation site, in particular 2-5 (additional) *in vivo* glycosylation sites, such as 2, 3, 4 or 5 (additional) *in vivo* glycosylation sites, preferably introduced by one or more of the substitutions described in any of the above lists.

Furthermore, the amino acid sequence of a polypeptide having at least one of the above mentioned *in vivo* N-glycosylation site modifications may differ from that of the parent polypeptide in that at least one cysteine residue has been introduced as identified above in the section entitled “*Conjugate of the invention having a non-polypeptide moiety attached to a cysteine residue*”, or at least one non-cysteine residue has been introduced as identified above in the section entitled “*Conjugate of the invention having a non-polypeptide moiety attaching to a non-cysteine residue*”.

Moreover, the polypeptide part of the conjugate of the invention may contain additional mutations, which are known to be advantageous. For example, in addition to the glycosylation sites discussed above, the polypeptide part of the conjugate may contain a substitution in a position selected from the group consisting of L194, A195, L228, Y249 and combinations thereof, in particular L194S, L194S+T245S and L194A+T254S (see WO 00/66754). Other examples of preferred additional substitutions include substitution or introduction of one or more glycosylation site(s) at or near positions known to be susceptible to proteolytic degradation. One position that is known to be susceptible to proteolytic degradation is H10 of wild-type human APC (see WO 98/48822).

It will be understood that in order to prepare a conjugate according to this aspect of the invention, the polypeptide must be expressed in a glycosylating host cell capable of attaching sugar moieties at the glycosylation sites or alternatively subjected to *in vitro* glycosylation. Examples of glycosylating host cells are given in the section below entitled “*Coupling to a sugar moiety*”.

Conjugate of the invention wherein the non-polypeptide moiety is attached to a cysteine residue

In another embodiment of the invention, the present invention relates to a conjugate comprising at least one non-polypeptide moiety, in particular a polymer molecule, covalently attached to a protein C polypeptide that comprises an amino acid sequence which differs from a parent protein C polypeptide, in particular from the amino acid sequence shown in SEQ ID NO:4 or a variant thereof, in at least one cysteine residue has been introduced and/or removed, in particular introduced. Thus, in an interesting embodiment of the invention the non-polypeptide moiety has cysteine as an attachment group. Preferably, the cysteine attachment group is introduced in a position which is occupied by an amino acid residue having at least 25% of its side chain exposed to the surface, such as at least 50% of its side chain exposed to the surface. Such amino acid residues are identified in Example 1 herein. Of particular interest among these positions are positions that in the parent polypeptide are occupied by a T or an S residue, preferably an S residue. In accordance herewith, an interesting cysteine-modified conjugate is one, wherein a cysteine residue has been introduced into at least one position selected from the group consisting of S3, S11, S12, T37, S42, S61, T68, S75, S77, S82, S99, S119, S153, S190, S216, S252, T253, T268, S270, S281, S304, S305, T315, S332, S336, S340, S367, and S416, and more preferably from the group consisting of S3, S11, S12, S42, S61, S75, S77, S82, S99, S119, S153, S190, S216, S252, S270, S281, S304, S305, S332, S336, S340, S367 and S416.

In a similar way as described above (see the section entitled “*Conjugate of the invention where the non-polypeptide moiety is a sugar moiety*” the cysteine residue is preferably introduced in a position which is within the active site region (defined in Example 2 herein) and which is occupied by an amino acid residue having at least 25% of its side chain exposed to the surface (defined in Example 3 herein), i.e. the cysteine residue is preferably introduced in a position selected from the group consisting of D172, D189, S190, K191, K192, K193, D214, E215, S216, K217, K218, L220, V243, V245, S250, K251, S252, T253, T254, D255, L296, Y302, H303, S304, S305, R306, E307, K308, E309, A310, R312, T315, F316, V334, S336, V339, M338, I348, L349, D351, R352, E357, G383, E385, L386, L387 and H388. More preferably, the cysteine residue is introduced in a positions selected from the group consisting of D189, S190, K191, D214, K217, K251, S252, T253, Y302, S305, E307, S336, V339, M338, G383 and L386.

The polypeptide part of the conjugate according to this embodiment typically comprises 1-10 introduced cysteine residues, in particular 1-5 or 1-3 introduced cysteine residues, e.g. 1, 2 or 3 introduced cysteine residues.

While the non-polypeptide moiety of the conjugate according to this aspect of the invention may be any molecule which, when using the given conjugation method has a cysteine residue as an attachment group (such as an polymer moiety, a lipophilic group or an organic derivatizing agent), it is preferred that the non-polypeptide moiety is a polymer molecule, e.g. any of the molecules mentioned in the section entitled "*Conjugation to a polymer molecule*". Preferably, the polymer molecule is selected from the group consisting of linear or branched polyethylene glycol or polyalkylene oxide. Most preferably, the polymer molecule is PEG, such as VS-PEG. The conjugation between the polypeptide and the polymer may be achieved in any suitable manner, e.g. as described in the section entitled "*Conjugation to a polymer molecule*", e.g. by using a one step method or by the stepwise manner referred to in said section. When the polypeptide comprises only one conjugatable cysteine residue, this is preferably conjugated to a first non-polypeptide moiety with a molecular weight of at least about 10kDa or at least about 15kDa, such as a molecular weight of about 12kDa, about 15kDa or about 20kDa, either directly conjugated or indirectly through a low molecular weight polymer (as disclosed in WO 99/55377). When the conjugate comprises two or more first non-polypeptide moieties, normally each of these has a molecular weight of about 5kDa, about 10kDa or about 12kDa.

The conjugate according to this embodiment may comprise at least one second non-polypeptide moiety, such as 1-10, 1-8, 1-5 or 1-3 such moieties. When the first non-polypeptide moiety is a polyalkylene oxide or PEG derived polymer, the second non-polypeptide moiety is preferably a sugar moiety, in particular an *in vivo* attached moiety. The sugar moiety may be present at one or more of the naturally-occurring glycosylation sites present in the parent polypeptide, or at an introduced glycosylation site. Suitable introduced glycosylation sites, in particular N-glycosylation sites, are described in the section entitled "*Conjugate of the invention wherein the non-polypeptide moiety is a sugar moiety*".

Moreover, the polypeptide part of the conjugate of the invention may contain additional mutations, which are known to be advantageous. For example, in addition to the introduced cysteine residues discussed above, the polypeptide part of the conjugate may contain a substitution in a position selected from the group consisting of L194, A195, L228, Y249 and combinations thereof, in particular L194S, L194S+T245S and L194A+T254S (see WO 00/66754). Other examples of preferred additional substitutions include substitution or introduction of one or more cysteine residue(s) at or near positions known to be susceptible to proteolytic degradation. One position that is known to be susceptible to proteolytic degradation is H10 of wild-type human APC (see WO 98/48822).

Conjugate of the invention wherein the non-polypeptide moiety is attached to a non-cysteine moiety

Based on the present disclosure the skilled person will be aware that amino acid residues comprising other attachment groups may be introduced by substitution into the parent polypeptide, using the same approach as that illustrated above with glycosylation sites and cysteine residues. For instance, one or more amino acid residues comprising an acid group (glutamic acid or aspartic acid), tyrosine, serine or lysine may be introduced into the positions discussed above (see the sections entitled “*Conjugate of the invention where the non-polypeptide moiety is a sugar moiety*” and “*Conjugate of the invention wherein the non-polypeptide moiety is attached to a cysteine residue*”).

Polypeptide variants of the invention

In a further aspect the present invention relates to generally novel variants of parent protein C polypeptides. The novel variants are important intermediate compounds for the preparation of conjugates of the invention. In addition, and as will be apparent from the below disclosure and from the examples provided herein, the variants themselves have interesting properties.

Thus, in its broadest aspect the present invention relates to novel variants of a parent protein C polypeptide, where the variants constitute the polypeptide part, more particularly the APC part, of the conjugates of the invention. As will be evident from the examples provided herein, it has been found that some variants, wherein one or more glycosylation sites were introduced, but not utilized, has interesting properties, in particular with respect to increased resistance towards inhibition by alpha-1-antitrypsin and increased resistance towards inactivation by human plasma. These variant comprises at least one substitution in the active site region (as defined in Example 2 herein), in particular they comprise a substitution of an amino acid residue, which is located in the active site region and which has at least 25% of its side chain exposed to the surface (as defined in Example 3 herein). Thus, preferred variants according to this aspect of the invention comprises a substitution in a position selected from the group consisting of D172, D189, S190, K191, K192, K193, D214, E215, S216, K217, K218, L220, V243, V245, S250, K251, S252, T253, T254, D255, L296, Y302, H303, S304, S305, R306, E307, K308, E309, A310, R312, T315, F316, V334, S336, N337, M338, I348, L349, D351, R352, E357, E382, G383, L386, L387 and H388, with the proviso that the substitution is not selected from

the group consisting of T254S, T254A, T254H, T254K, T254R, T254N, T254D, T254E, T254G, T254Q, Y302S, Y302A, Y302T, Y302H, Y302K, Y302R, Y302N, Y302D, Y302E, Y302G, Y302Q, F316S, F316A, F316T, F316H, F316K, F316R, F316N, F316D, F316E, F316G and F316Q.

5 As is evident from the above list of positions, which are located in the active site region and, at the same time, has at least 25% of its side chain exposed to the surface, a significant amount of the these positions are occupied by charged amino acid residues. Analysing the three-dimensional structure of protein C, in particular the above-identified region, it can be observed that at least some of the charged residues interact with each other. For example, K251 is
 10 believed to form a salt bridge to D214. Moreover, it can be seen that a cluster of negatively charged amino acid residues (D214, E215 and E357) is present. Without being bound by any particular theory it is contemplated that the charged amino acid residues within the above-identified region, or at least some of the charged amino acid residues within this particular region, are important for capturing and/or binding the substrate/inhibitor. Therefore, amino acid
 15 substitutions which are particular interesting according to this aspect of the present invention are constituted by such an amino acid substitutions, wherein a charged amino acid residue, which is located in the active site region and, at the same time, has at least 25% of its side chain exposed to the surface, is substituted with an amino acid residue having no charge, in particular an amino acid residue having no charge but a polar side chain (Gly, Ser, Thr, Cys, Tyr, Asn or
 20 Gln), as well as amino acid substitutions, wherein a charged amino acid residue, which is located in the active site region and, at the same time, has at least 25% of its side chain exposed to the surface, is substituted with an amino acid residue having an opposite charge.

Specific examples of amino acid substitutions, wherein the charge of the amino acid residue in question is changed to an opposite charge, include D172K, D172R, D189K, D189R,
 25 K191D, K191E, K192D, K192E, K193D, K193E, D214K, D214R, E215K, E215R, K217D, K217E, K218D, K218E, K251D, K251E, D255K, D255R, R306D, R306E, E307K, E307R, K308D, K308E, E309K, E309R, R312D, R312E, D351K, D351R, R352D, R352E, E357K, E357R, E382K and E382R, such as D214K, D214R, E215K, E215R, K251D, K251E, E357K and E357R, e.g. D214K, D214R, K251D and K251E.

30 Other specific examples of amino acid substitutions, wherein the charged amino acid residue in question is substituted with an amino acid side chain having a polar side chain, include D172G/S/T/C/Y/N/Q, D189G/S/T/C/Y/N/Q, K191G/S/T/C/Y/N/Q, K192G/S/T/C/Y/N/Q, K193G/S/T/C/Y/N/Q, D214G/S/T/C/Y/N/Q, E215G/S/T/C/Y/N/Q,

K217G/S/T/C/Y/N/Q, K218G/S/T/C/Y/N/Q, K251G/S/T/C/Y/N/Q, D255G/S/T/C/Y/N/Q, R306G/S/T/C/Y/N/Q, E307G/S/T/C/Y/N/Q, K308G/S/T/C/Y/N/Q, E309G/S/T/C/Y/N/Q, R312G/S/T/C/Y/N/Q, D351G/S/T/C/Y/N/Q, R352G/S/T/C/Y/N/Q, E357G/S/T/C/Y/N/Q and E382G/S/T/C/Y/N/Q, such as D214G/S/T/C/Y/N/Q, E215G/S/T/C/Y/N/Q,
5 K251G/S/T/C/Y/N/Q and E357G/S/T/C/Y/N/Q, e.g. D214Q, E215Q, K251Q and E357Q, in particular K251Q. Another interesting substitution may be K251N+T253A.

Further specific examples of interesting substitutions include the substitutions disclosed in the sections entitled "*Conjugate of the invention where the non-polypeptide moiety is a sugar moiety*" and "*Conjugate of the invention wherein the non-polypeptide moiety is at-*
10 *tached to a cysteine residue*", in particular the substitutions selected from the group consisting of K251N, S252N, Y302N and S190+K192T, especially K251N and S252N, most preferably K251N.

As will be understood, details and particulars concerning the conjugates of the invention (e.g. activation of protein C, number of substitutions, formulation of conjugates, indica-
15 tions for which the conjugates may be used, increased resistance towards inactivation by alpha-1-antitrypsin and human plasma, etc.) will be the same or analogous to the variant aspect of the invention, whenever appropriate. Thus, statements and details concerning the conjugates of the invention will apply mutatis mutandis to the protein C variants disclosed herein, whenever appropriate.

20

Non-polypeptide moiety of the conjugate of the invention

As indicated further above the non-polypeptide moiety of the conjugate of the invention is preferably selected from the group consisting of a polymer molecule, a lipophilic compound, a sugar moiety (by way of *in vivo* glycosylation) and an organic derivatizing agent. All
25 of these agents may confer desirable properties to the polypeptide part of the conjugate, in particular increased functional *in vivo* half-life and/or increased plasma half-life. The polypeptide part of the conjugate is normally conjugated to only one type of non-polypeptide moiety, but may also be conjugated to two or more different types of non-polypeptide moieties, e.g. to a polymer molecule and a sugar moiety, to a lipophilic group and a sugar moiety, to an organic
30 derivatizing agent and a sugar moiety, to a lipophilic group and a polymer molecule, etc. The conjugation to two or more different non-polypeptide moieties may be done simultaneous or sequentially.

Methods of preparing a conjugate of the invention

In the following sections “*Conjugation to a lipophilic compound*”, “*Conjugation to a polymer molecule*”, “*Conjugation to a sugar moiety*” and “*Conjugation to an organic derivatizing agent*” conjugation to specific types of non-polypeptide moieties is described. In general, a polypeptide conjugate according to the invention may be produced by culturing an appropriate host cell under conditions conducive for the expression of the polypeptide, and recovering the polypeptide, wherein a) the polypeptide comprises at least one N- or O-glycosylation site and the host cell is an eukaryotic host cell capable of *in vivo* glycosylation, and/or b) the polypeptide is subjected to conjugation to a non-polypeptide moiety *in vitro*.

It will be understood that the conjugation should be designed so as to produce the optimal molecule with respect to the number of non-polypeptide moieties attached, the size and form of such molecules (e.g. whether they are linear or branched), and the attachment site(s) in the polypeptide. The molecular weight of the non-polypeptide moiety to be used may e.g. be chosen on the basis of the desired effect to be achieved. For instance, if the primary purpose of the conjugation is to achieve a conjugate having a high molecular weight (e.g. to reduce renal clearance) it is usually desirable to conjugate as few high molecular weight non-polypeptide moieties as possible to obtain the desired molecular weight. When a high degree of shielding is desirable this may be obtained by use of a sufficiently high number of low molecular weight non-polypeptide moieties (e.g. with a molecular weight of from about 300 Da to about 5 kDa, such as a molecular weight of from 300 Da to 2 kDa).

Conjugation to a polymer molecule

The polymer molecule to be coupled to the polypeptide may be any suitable polymer molecule, such as a natural or synthetic homo-polymer or hetero-polymer, typically with a molecular weight in the range of about 300-100,000 Da, such as about 500-20,000 Da, more preferably in the range of about 500-15,000 Da, even more preferably in the range of about 2-12 kDa, such as in the range of about 3-10 kDa. When the term “about” is used herein in connection with a certain molecular weight, the word “about” indicates an approximate average molecular weight and reflects the fact that there will normally be a certain molecular weight distribution in a given polymer preparation.

Examples of homo-polymers include a polyol (i.e. poly-OH), a polyamine (i.e. poly-NH₂) and a polycarboxylic acid (i.e. poly-COOH). A hetero-polymer is a polymer comprising different coupling groups, such as a hydroxyl group and an amine group.

Examples of suitable polymer molecules include polymer molecules selected from the group consisting of polyalkylene oxide (PAO), including polyalkylene glycol (PAG), such as polyethylene glycol (PEG) and polypropylene glycol (PPG), branched PEGs, poly-vinyl alcohol (PVA), poly-carboxylate, poly-(vinylpyrrolidone), polyethylene-co-maleic acid anhydride, polystyrene-co-maleic acid anhydride, dextran, including carboxymethyl-dextran, or any other biopolymer suitable for reducing immunogenicity and/or increasing functional *in vivo* half-life and/or serum half-life. Another example of a polymer molecule is human albumin or another abundant plasma protein. Generally, polyalkylene glycol-derived polymers are biocompatible, non-toxic, non-antigenic, non-immunogenic, have various water solubility properties, and are easily excreted from living organisms.

PEG is the preferred polymer molecule, since it has only few reactive groups capable of cross-linking compared to, e.g., polysaccharides such as dextran. In particular, monofunctional PEG, e.g. methoxypolyethylene glycol (mPEG), is of interest since its coupling chemistry is relatively simple (only one reactive group is available for conjugating with attachment groups on the polypeptide). Consequently, the risk of cross-linking is eliminated, the resulting polypeptide conjugates are more homogeneous and the reaction of the polymer molecules with the polypeptide is easier to control.

To effect covalent attachment of the polymer molecule(s) to the polypeptide, the hydroxyl end groups of the polymer molecule must be provided in activated form, i.e. with reactive functional groups (examples of which include primary amino groups, hydrazide (HZ), thiol, succinate (SUC), succinimidyl succinate (SS), succinimidyl succinamide (SSA), succinimidyl propionate (SPA), succinimidyl butyrate (SBA), succinimidyl carboxymethylate (SCM), benzotriazole carbonate (BTC), N-hydroxysuccinimide (NHS), aldehyde, nitrophenylcarbonate (NPC), and tresylate (TRES)). Suitable activated polymer molecules are commercially available, e.g. from Shearwater Polymers, Inc., Huntsville, AL, USA, or from PolyMASC Pharmaceuticals plc, UK.

Alternatively, the polymer molecules can be activated by conventional methods known in the art, e.g. as disclosed in WO 90/13540. Specific examples of activated linear or branched polymer molecules for use in the present invention are described in the Shearwater Polymers, Inc. 1997 and 2000 Catalogs (Functionalized Biocompatible Polymers for Research and pharmaceuticals, Polyethylene Glycol and Derivatives, incorporated herein by reference).

Specific examples of activated PEG polymers include the following linear PEGs: NHS-PEG (e.g. SPA-PEG, SSPA-PEG, SBA-PEG, SS-PEG, SSA-PEG, SC-PEG, SG-PEG,

and SCM-PEG), and NOR-PEG, BTC-PEG, EPOX-PEG, NCO-PEG, NPC-PEG, CDI-PEG, ALD-PEG, TRES-PEG, VS-PEG, IODO-PEG and MAL-PEG, including the mPEG forms thereof, and branched PEGs such as PEG2-NHS, including the mPEG forms thereof, and those disclosed in US 5,932,462 and US 5,643,575, both of which are incorporated herein by reference. Furthermore, the following publications, incorporated herein by reference, disclose useful polymer molecules and/or PEGylation chemistries: US 5,824,778, US 5,476,653, WO 97/32607, EP 229,108, EP 402,378, US 4,902,502, US 5,281,698, US 5,122,614, US 5,219,564, WO 92/16555, WO 94/04193, WO 94/14758, WO 94/17039, WO 94/18247, WO 94/28024, WO 95/00162, WO 95/11924, WO95/13090, WO 95/33490, WO 96/00080, WO 97/18832, WO 98/41562, WO 98/48837, WO 99/32134, WO 99/32139, WO 99/32140, WO 96/40791, WO 98/32466, WO 95/06058, EP 439 508, WO 97/03106, WO 96/21469, WO 95/13312, EP 921 131, US 5,736,625, WO 98/05363, EP 809 996, US 5,629,384, WO 96/41813, WO 96/07670, US 5,473,034, US 5,516,673, EP 605 963, US 5,382,657, EP 510 356, EP 400 472, EP 183 503 and EP 154 316.

The conjugation of the polypeptide and the activated polymer molecules is conducted by use of any conventional method, e.g. as described in the following references (which also describe suitable methods for activation of polymer molecules): R.F. Taylor, (1991), "Protein immobilisation. Fundamental and applications", Marcel Dekker, N.Y.; S.S. Wong, (1992), "Chemistry of Protein Conjugation and Crosslinking", CRC Press, Florida, USA; G.T. Hermanson et al., (1993), "Immobilized Affinity Ligand Techniques", Academic Press, N.Y.). The skilled person will be aware that the activation method and/or conjugation chemistry to be used depends on the attachment group(s) of the polypeptide (examples of which are given further above), as well as the functional groups of the polymer (e.g. being amine, hydroxyl, carboxyl, aldehyde, sulfhydryl, succinimidyl, maleimide, vinylsulfone or haloacetate). The PEGylation may be directed towards conjugation to all available attachment groups on the polypeptide (i.e. such attachment groups that are exposed at the surface of the polypeptide) or may be directed towards one or more specific attachment groups, e.g. the N-terminal amino group as described in US 5,985,265. Furthermore, the conjugation may be achieved in one step or in a stepwise manner (e.g. as described in WO 99/55377).

It will be understood that the PEGylation is designed so as to produce the optimal molecule with respect to the number of PEG molecules attached, the size and form of such molecules (e.g. whether they are linear or branched), and the attachment site(s) in the polypep-

tide. The molecular weight of the polymer to be used may e.g. be chosen on the basis of the desired effect to be achieved.

In connection with conjugation to only a single attachment group on the protein (e.g. the N-terminal amino group), it may be advantageous that the polymer molecule, which may be linear or branched, has a high molecular weight, preferably about 10-25 kDa, such as about 15-25 kDa, e.g. about 20 kDa.

Normally, the polymer conjugation is performed under conditions aimed at reacting as many of the available polymer attachment groups with polymer molecules. This is achieved by means of a suitable molar excess of the polymer relative to the polypeptide. Typically, the molar ratios of activated polymer molecules to polypeptide are up to about 1000-1, such as up to about 200-1, or up to about 100-1. In some cases the ratio may be somewhat lower, however, such as up to about 50-1, 10-1, 5-1, 2-1 or 1-1 in order to obtain optimal reaction.

It is also contemplated according to the invention to couple the polymer molecules to the polypeptide through a linker. Suitable linkers are well known to the skilled person. A preferred example is cyanuric chloride (Abuchowski et al., (1977), J. Biol. Chem., 252, 3578-3581; US 4,179,337; Shafer et al., (1986), J. Polym. Sci. Polym. Chem. Ed., 24, 375-378).

Subsequent to the conjugation, residual activated polymer molecules are blocked according to methods known in the art, e.g. by addition of primary amine to the reaction mixture, and the resulting inactivated polymer molecules are removed by a suitable method.

It will be understood that depending on the circumstances, e.g. the amino acid sequence of the polypeptide, the nature of the activated PEG compound being used and the specific PEGylation conditions, including the molar ratio of PEG to polypeptide, varying degrees of PEGylation may be obtained, with a higher degree of PEGylation generally being obtained with a higher ratio of PEG to polypeptide. The PEGylated polypeptides resulting from any given PEGylation process will, however, normally comprise a stochastic distribution of polypeptide conjugates having slightly different degrees of PEGylation.

Coupling to a sugar moiety

In order to achieve *in vivo* glycosylation of a protein C molecule comprising one or more glycosylation sites the nucleotide sequence encoding the polypeptide must be inserted in a glycosylating, eucaryotic expression host. The expression host cell may be selected from fungal (filamentous fungal or yeast), insect or animal cells or from transgenic plant cells. In one embodiment the host cell is a mammalian cell, such as a COS cell, a CHO cell, a BHK cell or a

HEK cell, e.g. a HEK 293 cell, or an insect cell, such as an SF9 cell, or a yeast cell, e.g. *S. cerevisiae* or *Pichia pastoris*, or any of the host cells mentioned hereinafter.

Covalent *in vitro* coupling of sugar moieties (such as dextran) to amino acid residues of the polypeptide may also be used, e.g. as described, for example in WO 87/05330 and in
 5 Aplin et al., CRC Crit Rev. Biochem, pp. 259-306, 1981. The *in vitro* coupling of sugar moieties or PEG to protein- and peptide-bound Gln-residues can be carried out by transglutaminases (TGases). Transglutaminases catalyse the transfer of donor amine-groups to protein- and peptide-bound Gln-residues in a so-called cross-linking reaction. The donor-amine groups can be protein- or peptide-bound, such as the ϵ -amino-group in Lys-residues or it can be part of a
 10 small or large organic molecule. An example of a small organic molecule functioning as amino-donor in TGase-catalysed cross-linking is putrescine (1,4-diaminobutane). An example of a larger organic molecule functioning as amino-donor in TGase-catalysed cross-linking is an amine-containing PEG (Sato et al., 1996, Biochemistry 35, 13072-13080).

TGases, in general, are highly specific enzymes, and not every Gln-residues exposed
 15 on the surface of a protein is accessible to TGase-catalysed cross-linking to amino-containing substances. On the contrary, only few Gln-residues are naturally functioning as TGase substrates but the exact parameters governing which Gln-residues are good TGase substrates remain unknown. Thus, in order to render a protein susceptible to TGase-catalysed cross-linking reactions it is often a prerequisite at convenient positions to add stretches of amino acid sequence known to function very well as TGase substrates. Several amino acid sequences are
 20 known to be or to contain excellent natural TGase substrates e.g. substance P, elafin, fibrinogen, fibronectin, α_2 -plasmin inhibitor, α -caseins, and β -caseins.

Conjugation to an organic derivatizing agent

25 Covalent modification of the polypeptide may be performed by reacting one or more attachment groups of the polypeptide with an organic derivatizing agent. Suitable derivatizing agents and methods are well known in the art. For example, cysteinyl residues most commonly are reacted with α -haloacetates (and corresponding amines), such as chloroacetic acid or chloroacetamide, to give carboxymethyl or carboxyamidomethyl derivatives. Cysteinyl residues
 30 also are derivatized by reaction with bromotrifluoroacetone, α -bromo- β -(4-imidozoyl)propionic acid, chloroacetyl phosphate, N-alkylmaleimides, 3-nitro-2-pyridyl disulfide, methyl 2-pyridyl disulfide, p-chloromercuribenzoate, 2-chloromercuri-4-nitrophenol, or chloro-7-nitrobenzo-2-oxa-1,3-diazole. Histidyl residues are derivatized by reaction with diethylpyrocarbonate at pH

5.5-7.0 because this agent is relatively specific for the histidyl side chain. Para-bromophenacyl bromide also is useful. The reaction is preferably performed in 0.1 M sodium cacodylate at pH 6.0. Lysinyl and amino terminal residues are reacted with succinic or other carboxylic acid anhydrides. Derivatization with these agents has the effect of reversing the charge of the lysinyl residues. Other suitable reagents for derivatizing α -amino-containing residues include imidoesters such as methyl picolinimide, pyridoxal phosphate, pyridoxal, chloroborohydride, trinitrobenzenesulfonic acid, O-methylisourea, 2,4-pentanedione and transaminase-catalyzed reaction with glyoxylate. Arginyl residues are modified by reaction with one or several conventional reagents, among them phenylglyoxal, 2,3-butanedione, 1,2-cyclohexanedione, and ninhydrin. Derivatization of arginine residues requires that the reaction be performed in alkaline conditions because of the high pKa of the guanidine functional group.

Furthermore, these reagents may react with the groups of lysine as well as the arginine guanidino group. Carboxyl side groups (aspartyl or glutamyl) are selectively modified by reaction with carbodiimides ($R-N=C=N-R'$), where R and R' are different alkyl groups, such as 1-cyclohexyl-3-(2-morpholinyl-4-ethyl) carbodiimide or 1-ethyl-3-(4-azonia-4,4-dimethylpentyl) carbodiimide. Furthermore, aspartyl and glutamyl residues are converted to asparaginy and glutaminy residues by reaction with ammonium ions.

Conjugation to a lipophilic compound

The polypeptide and the lipophilic compound may be conjugated to each other, either directly or by use of a linker. The lipophilic compound may be a natural compound such as a saturated or unsaturated fatty acid, a fatty acid diketone, a terpene, a prostaglandin, a vitamin, a carotenoid or steroid, or a synthetic compound such as a carbon acid, an alcohol, an amine and sulphonic acid with one or more alkyl-, aryl-, alkenyl- or other multiple unsaturated compounds. The conjugation between the polypeptide and the lipophilic compound, optionally through a linker may be done according to methods known in the art, e.g. as described by Bodanszky in Peptide Synthesis, John Wiley, New York, 1976 and in WO 96/12505.

Conjugation of a tagged polypeptide

The polypeptide may be expressed as a fusion protein with a tag, i.e. an amino acid sequence or peptide stretch made up of typically 1-30, such as 1-20 amino acid residues. Besides allowing for fast and easy purification, the tag is a convenient tool for achieving conjugation between the tagged polypeptide and the non-polypeptide moiety. In particular, the tag may

be used for achieving conjugation in microtiter plates or other carriers, such as paramagnetic beads, to which the tagged polypeptide can be immobilised via the tag. The conjugation to the tagged polypeptide in, e.g., microtiter plates has the advantage that the tagged polypeptide can be immobilised in the microtiter plates directly from the culture broth (in principle without any purification) and subjected to conjugation. Thereby, the total number of process steps (from expression to conjugation) can be reduced. Furthermore, the tag may function as a spacer molecule, ensuring an improved accessibility to the immobilised polypeptide to be conjugated. The conjugation using a tagged polypeptide may be to any of the non-polypeptide moieties disclosed herein, e.g. to a polymer molecule such as PEG.

The identity of the specific tag to be used is not critical as long as the tag is capable of being expressed with the polypeptide and is capable of being immobilised on a suitable surface or carrier material. A number of suitable tags are commercially available, e.g. from Unizyme Laboratories, Denmark. For instance, the tag may consist of any of the following sequences:

His-His-His-His-His-His

Met-Lys-His-His-His-His-His

Met-Lys-His-His-Ala-His-His-Gln-His-His

Met-Lys-His-Gln-His-Gln-His-Gln-His-Gln-His-Gln

Met-Lys-His-Gln-His-Gln-His-Gln-His-Gln-His-Gln-Gln

or any of the following:

EQKLI SEEDL (a C-terminal tag described in Mol. Cell. Biol. 5:3610-16, 1985)

DYKDDDDK (a C- or N-terminal tag)

YPYDVPDYA

Antibodies against the above tags are commercially available, e.g. from ADI, Aves Lab and Research Diagnostics.

The subsequent cleavage of the tag from the polypeptide may be achieved by use of commercially available enzymes.

Methods of preparing a polypeptide variant of the invention or the polypeptide part of the conjugate of the invention

The polypeptide variant of the present invention or the polypeptide part of a conjugate of the invention, optionally in glycosylated form, may be produced by any suitable method known in the art. Such methods include constructing a nucleotide sequence encoding the polypeptide and expressing the sequence in a suitable transformed or transfected host. Preferably, the host cell is a gammacarboxylating host cell such as a mammalian cell. However,

the host cell is a gammacarboxylating host cell such as a mammalian cell. However, polypeptides of the invention may be produced, albeit less efficiently, by chemical synthesis or a combination of chemical synthesis or a combination of chemical synthesis and recombinant DNA technology.

5 A nucleotide sequence encoding a polypeptide variant or the polypeptide part of a conjugate of the invention may be constructed by isolating or synthesizing a nucleotide sequence encoding the parent protein C, such as protein C with the amino acid sequence shown in SEQ ID NO:2 and 4 and then changing the nucleotide sequence so as to effect introduction (i.e. insertion or substitution) or removal (i.e. deletion or substitution) of the relevant amino acid residue(s).

The nucleotide sequence is conveniently modified by site-directed mutagenesis in accordance with conventional methods. Alternatively, the nucleotide sequence is prepared by chemical synthesis, e.g. by using an oligonucleotide synthesizer, wherein oligonucleotides are designed based on the amino acid sequence of the desired polypeptide, and preferably selecting
15 those codons that are favored in the host cell in which the recombinant polypeptide will be produced. For example, several small oligonucleotides coding for portions of the desired polypeptide may be synthesized and assembled by PCR, ligation or ligation chain reaction (LCR) (Barany, PNAS 88:189-193, 1991). The individual oligonucleotides typically contain 5' or 3' overhangs for complementary assembly.

20 Alternative nucleotide sequence modification methods are available for producing polypeptide variants for high throughput screening, for instance methods which involve homologous cross-over such as disclosed in US 5,093,257, and methods which involve gene shuffling, i.e. recombination between two or more homologous nucleotide sequences resulting in new nucleotide sequences having a number of nucleotide alterations when compared to the
25 starting nucleotide sequences. Gene shuffling (also known as DNA shuffling) involves one or more cycles of random fragmentation and reassembly of the nucleotide sequences, followed by screening to select nucleotide sequences encoding polypeptides with desired properties. In order for homology-based nucleic acid shuffling to take place, the relevant parts of the nucleotide sequences are preferably at least 50% identical, such as at least 60% identical, more preferably
30 at least 70% identical, such as at least 80% identical. The recombination can be performed *in vitro* or *in vivo*.

Examples of suitable *in vitro* gene shuffling methods are disclosed by Stemmer et al. (1994), Proc. Natl. Acad. Sci. USA; vol. 91, pp. 10747-10751; Stemmer (1994), Nature, vol.

370, pp. 389-391; Smith (1994), Nature vol. 370, pp. 324-325; Zhao et al., Nat. Biotechnol. 1998, Mar; 16(3): 258-61; Zhao H. and Arnold, FB, Nucleic Acids Research, 1997, Vol. 25. No. 6 pp. 1307-1308; Shao et al., Nucleic Acids Research 1998, Jan 15; 26(2): pp. 681-83; and WO 95/17413.

5 An example of a suitable *in vivo* shuffling method is disclosed in WO 97/07205. Other techniques for mutagenesis of nucleic acid sequences by *in vitro* or *in vivo* recombination are disclosed e.g. in WO 97/20078 and US 5,837,458. Examples of specific shuffling techniques include "family shuffling", "synthetic shuffling" and "*in silico* shuffling".

10 Family shuffling involves subjecting a family of homologous genes from different species to one or more cycles of shuffling and subsequent screening or selection. Family shuffling techniques are disclosed e.g. by Crameri et al. (1998), Nature, vol. 391, pp. 288-291; Christians et al. (1999), Nature Biotechnology, vol. 17, pp. 259-264; Chang et al. (1999), Nature Biotechnology, vol. 17, pp. 793-797; and Ness et al. (1999), Nature Biotechnology, vol. 17, 893-896.

15 Synthetic shuffling involves providing libraries of overlapping synthetic oligonucleotides based e.g. on a sequence alignment of homologous genes of interest. The synthetically generated oligonucleotides are recombined, and the resulting recombinant nucleic acid sequences are screened and if desired used for further shuffling cycles. Synthetic shuffling techniques are disclosed in WO 00/42561.

20 *In silico* shuffling refers to a DNA shuffling procedure, which is performed or modelled using a computer system, thereby partly or entirely avoiding the need for physically manipulating nucleic acids. Techniques for *in silico* shuffling are disclosed in WO 00/42560. Once assembled (by synthesis, site-directed mutagenesis or another method), the nucleotide sequence encoding the polypeptide is inserted into a recombinant vector and operably linked to
25 control sequences necessary for expression of protein C in the desired transformed host cell.

It should of course be understood that not all vectors and expression control sequences function equally well to express the nucleotide sequence encoding a polypeptide described herein. Neither will all hosts function equally well with the same expression system. However, one of skill in the art may make a selection among these vectors, expression control sequences
30 and hosts without undue experimentation. For example, in selecting a vector, the host must be considered because the vector must replicate in it or be able to integrate into the chromosome. The vector's copy number, the ability to control that copy number, and the expression of any other proteins encoded by the vector, such as antibiotic markers, should also be considered. In

selecting an expression control sequence, a variety of factors should also be considered. These include, for example, the relative strength of the sequence, its controllability, and its compatibility with the nucleotide sequence encoding the polypeptide, particularly as regards potential secondary structures. Hosts should be selected by consideration of their compatibility with the chosen vector, the toxicity of the product coded for by the nucleotide sequence, their secretion characteristics, their ability to fold the polypeptide correctly, their fermentation or culture requirements, and the ease of purification of the products coded for by the nucleotide sequence. The recombinant vector may be an autonomously replicating vector, i.e. a vector, which exists as an extrachromosomal entity, the replication of which is independent of chromosomal replication, e.g. a plasmid. Alternatively, the vector is one which, when introduced into a host cell, is integrated into the host cell genome and replicated together with the chromosome(s) into which it has been integrated.

The vector is preferably an expression vector, in which the nucleotide sequence encoding the polypeptide of the invention is operably linked to additional segments required for transcription of the nucleotide sequence. The vector is typically derived from plasmid or viral DNA. A number of suitable expression vectors for expression in the host cells mentioned herein are commercially available or described in the literature. Useful expression vectors for eukaryotic hosts, include, for example, vectors comprising expression control sequences from SV40, bovine papilloma virus, adenovirus and cytomegalovirus. Specific vectors are, e.g., pCDNA3.1(+)\Hyg (Invitrogen, Carlsbad, CA, USA) and pCI-neo (Stratagene, La Jolla, CA, USA). Useful expression vectors for yeast cells include the 2 μ plasmid and derivatives thereof, the POT1 vector (US 4,931,373), the pJSO37 vector described in Okkels, Ann. New York Acad. Sci. 782, 202-207, 1996, and pPICZ A, B or C (Invitrogen). Useful vectors for insect cells include pVL941, pBG311 (Cate et al., "Isolation of the Bovine and Human Genes for Mullerian Inhibiting Substance And Expression of the Human Gene In Animal Cells", Cell, 45, pp. 685-98 (1986), pBluebac 4.5 and pMelbac (both available from Invitrogen). Useful expression vectors for bacterial hosts include known bacterial plasmids, such as plasmids from E. coli, including pBR322, pET3a and pET12a (both from Novagen Inc., WI, USA), wider host range plasmids, such as RP4, phage DNAs, e.g., the numerous derivatives of phage lambda, e.g., NM989, and other DNA phages, such as M13 and filamentous single stranded DNA phages.

Other vectors for use in this invention include those that allow the nucleotide sequence encoding the polypeptide to be amplified in copy number. Such amplifiable vectors are well known in the art. They include, for example, vectors able to be amplified by DHFR amplifica-

tion (see, e.g., Kaufman, U.S. Pat. No. 4,470,461, Kaufman and Sharp, "Construction Of A Modular Dihydrofolate Reductase cDNA Gene: Analysis Of Signals Utilized For Efficient Expression", Mol. Cell. Biol., 2, pp. 1304-19 (1982)) and glutamine synthetase ("GS") amplification (see, e.g., US 5,122,464 and EP 338,841).

5 The recombinant vector may further comprise a DNA sequence enabling the vector to replicate in the host cell in question. An example of such a sequence (when the host cell is a mammalian cell) is the SV40 origin of replication. When the host cell is a yeast cell, suitable sequences enabling the vector to replicate are the yeast plasmid 2 μ replication genes REP 1-3 and origin of replication.

10 The vector may also comprise a selectable marker, e.g. a gene the product of which complements a defect in the host cell, such as the gene coding for dihydrofolate reductase (DHFR) or the Schizosaccharomyces pombe TPI gene (described by P.R. Russell, Gene 40, 1985, pp. 125-130), or one which confers resistance to a drug, e.g. ampicillin, kanamycin, tetracyclin, chloramphenicol, neomycin, hygromycin or methotrexate. For Saccharomyces cere-
15 visiae, selectable markers include ura3 and leu2. For filamentous fungi, selectable markers include amdS, pyrG, arcB, niaD and sC.

 The term "control sequences" is defined herein to include all components, which are necessary or advantageous for the expression of the polypeptide of the invention. Each control sequence may be native or foreign to the nucleic acid sequence encoding the polypeptide. Such
20 control sequences include, but are not limited to, a leader sequence, polyadenylation sequence, propeptide sequence, promoter, enhancer or upstream activating sequence, signal peptide sequence, and transcription terminator. At a minimum, the control sequences include a promoter.

 A wide variety of expression control sequences may be used in the present invention. Such useful expression control sequences include the expression control sequences associated
25 with structural genes of the foregoing expression vectors as well as any sequence known to control the expression of genes of prokaryotic or eukaryotic cells or their viruses, and various combinations thereof.

 Examples of suitable control sequences for directing transcription in mammalian cells include the early and late promoters of SV40 and adenovirus, e.g. the adenovirus 2 major late
30 promoter, the MT-1 (metallothionein gene) promoter, the human cytomegalovirus immediate-early gene promoter (CMV), the human elongation factor 1 α (EF-1 α) promoter, the Drosophila minimal heat shock protein 70 promoter, the Rous Sarcoma Virus (RSV) promoter, the human ubiquitin C (UbC) promoter, the human growth hormone terminator, SV40 or adenovirus Elb

region polyadenylation signals and the Kozak consensus sequence (Kozak, M. J Mol Biol 1987 Aug 20;196(4):947-50).

In order to improve expression in mammalian cells a synthetic intron may be inserted in the 5' untranslated region of the nucleotide sequence encoding the polypeptide. An example
5 of a synthetic intron is the synthetic intron from the plasmid pCI-Neo (available from Promega Corporation, WI, USA).

Examples of suitable control sequences for directing transcription in insect cells include the polyhedrin promoter, the P10 promoter, the *Autographa californica* polyhedrosis virus basic protein promoter, the baculovirus immediate early gene 1 promoter and the baculovirus
10 39K delayed-early gene promoter, and the SV40 polyadenylation sequence. Examples of suitable control sequences for use in yeast host cells include the promoters of the yeast α -mating system, the yeast triose phosphate isomerase (TPI) promoter, promoters from yeast glycolytic genes or alcohol dehydrogenase genes, the ADH2-4c promoter, and the inducible GAL promoter. Examples of suitable control sequences for use in filamentous fungal host cells include
15 the ADH3 promoter and terminator, a promoter derived from the genes encoding *Aspergillus oryzae* TAKA amylase triose phosphate isomerase or alkaline protease, an *A. niger* α -amylase, *A. niger* or *A. nidulans* glucoamylase, *A. nidulans* acetamidase, *Rhizomucor miehei* aspartic proteinase or lipase, the TPI1 terminator and the ADH3 terminator. Examples of suitable control sequences for use in bacterial host cells include promoters of the lac system, the trp system,
20 the TAC or TRC system, and the major promoter regions of phage lambda.

The presence or absence of a signal peptide will, e.g., depend on the expression host cell used for the production of the polypeptide to be expressed (whether it is an intracellular or extracellular polypeptide) and whether it is desirable to obtain secretion. For use in filamentous fungi, the signal peptide may conveniently be derived from a gene encoding an *Aspergillus* sp.
25 amylase or glucoamylase, a gene encoding a *Rhizomucor miehei* lipase or protease or a *Humicola lanuginosa* lipase. The signal peptide is preferably derived from a gene encoding *A. oryzae* TAKA amylase, *A. niger* neutral α -amylase, *A. niger* acid-stable amylase, or *A. niger* glucoamylase. For use in insect cells, the signal peptide may conveniently be derived from an insect gene (cf. WO 90/05783), such as the Lepidopteran *manduca sexta* adipokinetic hormone precursor, (cf. US 5,023,328), the honeybee melittin (Invitrogen), ecdysteroid UDPglucosyltransferase (egt) (Murphy et al., Protein Expression and Purification 4, 349-357 (1993) or human
30 pancreatic lipase (hpl) (Methods in Enzymology 284, pp. 262-272, 1997). A preferred signal peptide for use in mammalian cells is that of hFVII or the murine Ig kappa light chain signal

peptide (Coloma, M (1992) J. Imm. Methods 152:89-104). For use in yeast cells suitable signal peptides have been found to be the α -factor signal peptide from *S. cerevisiae* (cf. US 4,870,008), a modified carboxypeptidase signal peptide (cf. L.A. Valls et al., Cell 48, 1987, pp. 887-897), the yeast BAR1 signal peptide (cf. WO 87/02670), the yeast aspartic protease 3 (YAP3) signal peptide (cf. M. Egel-Mitani et al., Yeast 6, 1990, pp. 127-137), and the synthetic leader sequence TA57 (WO98/32867). For use in *E. coli* cells a suitable signal peptide have been found to be the signal peptide ompA (EP581821).

The nucleotide sequence of the invention encoding a protein C polypeptide variant, whether prepared by site-directed mutagenesis, synthesis, PCR or other methods, may optionally include a nucleotide sequence that encode a signal peptide. The signal peptide is present when the polypeptide is to be secreted from the cells in which it is expressed. Such signal peptide, if present, should be one recognized by the cell chosen for expression of the polypeptide. The signal peptide may be homologous (e.g. be that normally associated with human protein C) or heterologous (i.e. originating from another source than human protein C) to the polypeptide or may be homologous or heterologous to the host cell, i.e. be a signal peptide normally expressed from the host cell or one which is not normally expressed from the host cell. Accordingly, the signal peptide may be prokaryotic, e.g. derived from a bacterium such as *E. coli*, or eukaryotic, e.g. derived from a mammalian, or insect or yeast cell.

Any suitable host may be used to produce the polypeptide or polypeptide part of the conjugate of the invention, including bacteria, fungi (including yeasts), plant, insect, mammal, or other appropriate animal cells or cell lines, as well as transgenic animals or plants. Examples of bacterial host cells include grampositive bacteria such as strains of *Bacillus*, e.g. *B. brevis* or *B. subtilis*, *Pseudomonas* or *Streptomyces*, or gramnegative bacteria, such as strains of *E. coli*. The introduction of a vector into a bacterial host cell may, for instance, be effected by protoplast transformation (see, e.g., Chang and Cohen, 1979, Molecular General Genetics 168: 111-115), using competent cells (see, e.g., Young and Spizizin, 1961, Journal of Bacteriology 81: 823-829, or Dubnau and Davidoff-Abelson, 1971, Journal of Molecular Biology 56: 209-221), electroporation (see, e.g., Shigekawa and Dower, 1988, Biotechniques 6: 742-751), or conjugation (see, e.g., Koehler and Thorne, 1987, Journal of Bacteriology 169: 5771-5278). Examples of suitable filamentous fungal host cells include strains of *Aspergillus*, e.g. *A. oryzae*, *A. niger*, or *A. nidulans*, *Fusarium* or *Trichoderma*. Fungal cells may be transformed by a process involving protoplast formation, transformation of the protoplasts, and regeneration of the cell wall in a manner known per se. Suitable procedures for transformation of *Aspergillus* host cells are de-

scribed in EP 238 023 and US 5,679,543. Suitable methods for transforming *Fusarium* species are described by Malardier et al., 1989, Gene 78: 147-156 and WO 96/00787. Examples of suitable yeast host cells include strains of *Saccharomyces*, e.g. *S. cerevisiae*, *Schizosaccharomyces*, *Kluyveromyces*, *Pichia*, such as *P. pastoris* or *P. methanolica*, *Hansenula*, such as *H. Polymorpha* or *Yarrowia*. Yeast may be transformed using the procedures described by Becker and Guar-
5 ente, In Abelson, J.N. and Simon, M.I., editors, Guide to Yeast Genetics and Molecular Biology, Methods in Enzymology, Volume 194, pp 182-187, Academic Press, Inc., New York; Ito et al., 1983, Journal of Bacteriology 153: 163; Hinnen et al., 1978, Proceedings of the National Academy of Sciences USA 75: 1920; and as disclosed by Clontech Laboratories, Inc, Palo Alto,
10 CA, USA (in the product protocol for the YeastmakerTM Yeast Transformation System Kit). Examples of suitable insect host cells include a *Lepidoptera* cell line, such as *Spodoptera frugiperda* (Sf9 or Sf21) or *Trichoplusia ni* cells (High Five) (US 5,077,214). Transformation of insect cells and production of heterologous polypeptides therein may be performed as described by Invitrogen. Examples of suitable mammalian host cells include Chinese hamster
15 ovary (CHO) cell lines, (e.g. CHO-K1; ATCC CCL-61), Green Monkey cell lines (COS) (e.g. COS 1 (ATCC CRL-1650), COS 7 (ATCC CRL-1651)); mouse cells (e.g. NS/O), Baby Hamster Kidney (BHK) cell lines (e.g. ATCC CRL-1632 or ATCC CCL-10), and human cells (e.g. HEK 293 (ATCC CRL-1573)), as well as plant cells in tissue culture. Additional suitable cell
20 lines are known in the art and available from public depositories such as the American Type Culture Collection, Rockville, Maryland. Also, the mammalian cell, such as a CHO cell, may be modified to express sialyltransferase, e.g. 1,6-sialyltransferase, e.g. as described in US 5,047,335, in order to provide improved glycosylation of the protein C polypeptide.

In order to increase secretion it may be of particular interest to produce the polypeptide of the invention together with an endoprotease, in particular a PACE (Paired basic amino acid
25 converting enzyme) (e.g. as described in US 5,986,079), such as a Kex2 endoprotease (e.g. as described in WO 00/28065).

Methods for introducing exogenous DNA into mammalian host cells include calcium phosphate-mediated transfection, electroporation, DEAE-dextran mediated transfection, liposome-mediated transfection, viral vectors and the transfection method described by Life Technologies Ltd, Paisley, UK using Lipofectamin 2000. These methods are well known in the art
30 and e.g. described by Ausbel et al. (eds.), 1996, Current Protocols in Molecular Biology, John Wiley & Sons, New York, USA. The cultivation of mammalian cells are conducted according to established methods, e.g. as disclosed in (Animal Cell Biotechnology, Methods and Proto-

cols, Edited by Nigel Jenkins, 1999, Human Press Inc, Totowa, New Jersey, USA and Harrison MA and Rae IF, General Techniques of Cell Culture, Cambridge University Press 1997).

In the production methods of the present invention, the cells are cultivated in a nutrient medium suitable for production of the polypeptide using methods known in the art. For example, the cell may be cultivated by shake flask cultivation, small-scale or large-scale fermentation (including continuous, batch, fed-batch, or solid state fermentations) in laboratory or industrial fermenters performed in a suitable medium and under conditions allowing the polypeptide to be expressed and/or isolated. The cultivation takes place in a suitable nutrient medium comprising carbon and nitrogen sources and inorganic salts, using procedures known in the art.

Suitable media are available from commercial suppliers or may be prepared according to published compositions (e.g., in catalogues of the American Type Culture Collection). If the polypeptide is secreted into the nutrient medium, the polypeptide can be recovered directly from the medium. If the polypeptide is not secreted, it can be recovered from cell lysates.

The resulting polypeptide may be recovered by methods known in the art. For example, the polypeptide may be recovered from the nutrient medium by conventional procedures including, but not limited to, centrifugation, filtration, ultra-filtration, extraction or precipitation.

The polypeptides may be purified by a variety of procedures known in the art including, but not limited to, chromatography (e.g., ion exchange, affinity, hydrophobic, chromatofocusing, and size exclusion), electrophoretic procedures (e.g., preparative isoelectric focusing), differential solubility (e.g., ammonium sulfate precipitation) or extraction (see, e.g., Protein Purification, J.-C. Janson and Lars Ryden, editors, VCH Publishers, New York, 1989).

Pharmaceutical compositions and use

In a further aspect, the present invention relates to a pharmaceutical composition comprising a conjugate of the invention or a variant of the invention and a pharmaceutically acceptable carrier or excipient. In the present context, the term "Pharmaceutically acceptable" means that the carrier or excipient, at the dosages and concentrations employed, will not cause any unwanted or harmful effects in the patients to which they are administered. Such pharmaceutically acceptable carriers and excipients are well known in the art (see Remington's Pharmaceutical Sciences, 18th edition, A. R. Gennaro, Ed., Mack Publishing Company [1990]; Pharmaceutical Formulation Development of Peptides and Proteins, S. Frokjaer and L. Hovgaard, Eds.,

Taylor & Francis [2000] ; and Handbook of Pharmaceutical Excipients, 3rd edition, A. Kibbe, Ed., Pharmaceutical Press [2000]).

In a still further aspect, the present invention relates to a conjugate of the invention, a variant of the invention or a pharmaceutical composition of the invention for use as a medica-
5 ment. More particularly, the conjugates, variants or pharmaceutical compositions of the invention may be used for the manufacture of a medicament for the treatment of stroke; myocardial infarction; after venous thrombosis; disseminated intravascular coagulation (DIC); sepsis; septic shock; emboli, such as pulmonary emboli; transplantation, such as bone marrow transplanta-
tion; burns; pregnancy; major surgery/traum or adult respiratory stress syndrome (ARDS), in
10 particular for the treatment of septic shock

The present invention also relates to a method for treating or preventing a disease selected from the group consisting of stroke; myocardial infarction; after venous thrombosis; disseminated intravascular coagulation (DIC); sepsis; septic shock; emboli, such as pulmonary
emboli; transplantation, such as bone marrow transplantation; burns; pregnancy; major sur-
15 gery/traum and adult respiratory stress syndrome (ARDS), the method comprising administering to a patient in need thereof an effective amount of a conjugate of the invention, of a variant according to the invention, or of a pharmaceutical composition according to the invention, in particular for treating or preventing, especially treating, septic shock.

A "patient" for the purposes of the present invention includes both humans and other
20 mammals. Thus the methods are applicable to both human therapy and veterinary applications.

The polypeptide variants and conjugates of the invention will be administered to patients in an effective dose. By "effective dose" herein is meant a dose that is sufficient to produce the desired effects in relation to the condition for which it is administered. The exact dose will depend on the disorder to be treated, and will be ascertainable by one skilled in the art using
25 known techniques. As mentioned above, in the treatment of severe sepsis 24 µg/kg/h of human APC is administered for 96 hours, which corresponds to a total amount of protein of about 230 mg for a patient having a body weight of about 100 kg. The conjugates and variants of the present invention are, due to their increased plasma half-lives, contemplated to have a higher efficacy due to the extended action-time in plasma. This increased efficacy may, for ex-
30 ample, be estimated by calculating the area under the curve (AUC) in the "Human Plasma Inactivation assay II" or by measuring the serum half-life. The increased efficacy means that the effective dose needed to obtain the desired effect for a particular disorder will be smaller (less protein need to be administered) than the effective dose of human APC. In addition, the in-

creased plasma half-life will also allow treatment where the APC variants or conjugates are used regularly with a given time-period. Thus, these new properties will permit the use of a reduced amount and/or and less frequent administration, such as bolus injections, of the compounds of the invention. For example, the compounds of the invention may be administered by
5 a either a bolus or infusion or as a combination thereof with doses which range from 1 $\mu\text{g/kg}$ body weight as a bolus every 2nd hour for several days (e.g. for 96 hours) to 1 mg/kg body weight as a bolus once every 4th day. Preferably, as low a dose as possible is administered as less frequent as possible, e.g. 1-500 $\mu\text{g/kg}$ body weight, preferably 1-250 $\mu\text{g/kg}$ body weight, such as 1-100 $\mu\text{g/kg}$ body weight, more preferably 1-50 $\mu\text{g/kg}$ body weight is administered as a
10 bolus every 4-96 hour, e.g. every 8-96 hour, such as every 16-96, every 24-96 hour, every 40-96 hour, every 48-96 hour, every 56-96 hour, every 72-96 hour.

Compounds of the invention, which are preferred are such compounds where the ratio between the AUC of said compound, in its activated form, and the AUC of human APC is at least 1.25 when tested in the "Human Plasma Inactivation Assay II" described in Example 13
15 herein. Preferably, the ratio is at least 1.5, such as at least 2, e.g. at least 3, more preferably the ratio is at least 4, such as at least 5, e.g. at least 6, even more preferably the ratio is at least 7, such as at least 8, e.g. at least 9, most preferably the ratio is at least 10.

The polypeptide variant or conjugate of the invention can be used "as is" and/or in a salt form thereof. Suitable salts include, but are not limited to, salts with alkali metals or alkali
20 line earth metals, such as sodium, potassium, calcium and magnesium, as well as e.g. zinc salts. These salts or complexes may be present as a crystalline and/or amorphous structure.

The pharmaceutical composition of the invention may be administered alone or in conjunction with other therapeutic agents. These agents may be incorporated as part of the same pharmaceutical composition or may be administered separately from the polypeptide or conjugate
25 of the invention, either concurrently or in accordance with another treatment schedule. In addition, the polypeptide, conjugate or pharmaceutical composition of the invention may be used as an adjuvant to other therapies.

The pharmaceutical composition of the invention may be formulated in a variety of forms, e.g. as a liquid, gel, lyophilized, or as a compressed solid. The preferred form will depend upon the particular indication being treated and will be readily able to be determined by
30 one skilled in the art.

The administration of the formulations of the present invention can be performed in a variety of ways, including, but not limited to, orally, subcutaneously, intravenously, intracere-

brally, intranasally, transdermally, intraperitoneally, intramuscularly, intrapulmonary, vaginally, rectally, intraocularly, or in any other acceptable manner. The formulations can be administered continuously by infusion, although bolus injection is acceptable, using techniques well known in the art, such as pumps or implantation. In some instances the formulations may
5 be directly applied as a solution or spray.

Parenteral compositions

An example of a pharmaceutical composition is a solution designed for parenteral administration. Although in many cases pharmaceutical solution formulations are provided in
10 liquid form, appropriate for immediate use, such parenteral formulations may also be provided in frozen or in lyophilized form. In the former case, the composition must be thawed prior to use. The latter form is often used to enhance the stability of the active compound contained in the composition under a wider variety of storage conditions, as it is recognized by those skilled in the art that lyophilized preparations are generally more stable than their liquid counterparts.
15 Such lyophilized preparations are reconstituted prior to use by the addition of one or more suitable pharmaceutically acceptable diluents such as sterile water for injection or sterile physiological saline solution.

In case of parenterals, they are prepared for storage as lyophilized formulations or aqueous solutions by mixing, as appropriate, the polypeptide having the desired degree of purity with one or more pharmaceutically acceptable carriers, excipients or stabilizers typically
20 employed in the art (all of which are termed "excipients"), for example buffering agents, stabilizing agents, preservatives, isotonifiers, non-ionic detergents, antioxidants and/or other miscellaneous additives.

Buffering agents help to maintain the pH in the range which approximates physiological conditions. They are typically present at a concentration ranging from about 2 mM to about
25 50 mM Suitable buffering agents for use with the present invention include both organic and inorganic acids and salts thereof such as citrate buffers (e.g., monosodium citrate-disodium citrate mixture, citric acid-trisodium citrate mixture, citric acid-monosodium citrate mixture, etc.), succinate buffers (e.g., succinic acid-monosodium succinate mixture, succinic acid-sodium hydroxide mixture, succinic acid-disodium succinate mixture, etc.), tartrate buffers (e.g., tartaric
30 acid-sodium tartrate mixture, tartaric acid-potassium tartrate mixture, tartaric acid-sodium hydroxide mixture, etc.), fumarate buffers (e.g., fumaric acid-monosodium fumarate mixture, fumaric acid-disodium fumarate mixture, monosodium fumarate-disodium fumarate mixture,

etc.), gluconate buffers (e.g., gluconic acid-sodium glyconate mixture, gluconic acid-sodium hydroxide mixture, gluconic acid-potassium glyuconate mixture, etc.), oxalate buffer (e.g., oxalic acid-sodium oxalate mixture, oxalic acid-sodium hydroxide mixture, oxalic acid-potassium oxalate mixture, etc.), lactate buffers (e.g., lactic acid-sodium lactate mixture, lactic acid-sodium hydroxide mixture, lactic acid-potassium lactate mixture, etc.) and acetate buffers (e.g., acetic acid-sodium acetate mixture, acetic acid-sodium hydroxide mixture, etc.). Additional possibilities are phosphate buffers, histidine buffers and trimethylamine salts such as Tris. Preservatives are added to retard microbial growth, and are typically added in amounts of e.g. about 0.1%-2% (w/v). Suitable preservatives for use with the present invention include phenol, benzyl alcohol, meta-cresol, methyl paraben, propyl paraben, octadecyldimethylbenzyl ammonium chloride, benzalkonium halides (e.g. benzalkonium chloride, bromide or iodide), hexamethonium chloride, alkyl parabens such as methyl or propyl paraben, catechol, resorcinol, cyclohexanol and 3-pentanol.

Isotonicifiers are added to ensure isotonicity of liquid compositions and include polyhydric sugar alcohols, preferably trihydric or higher sugar alcohols, such as glycerin, erythritol, arabitol, xylitol, sorbitol and mannitol. Polyhydric alcohols can be present in an amount between 0.1% and 25% by weight, typically 1% to 5%, taking into account the relative amounts of the other ingredients.

Stabilizers refer to a broad category of excipients which can range in function from a bulking agent to an additive which solubilizes the therapeutic agent or helps to prevent denaturation or adherence to the container wall. Typical stabilizers can be polyhydric sugar alcohols (enumerated above); amino acids such as arginine, lysine, glycine, glutamine, asparagine, histidine, alanine, omithine, L-leucine, 2-phenylalanine, glutamic acid, threonine, etc., organic sugars or sugar alcohols, such as lactose, trehalose, stachyose, mannitol, sorbitol, xylitol, ribitol, myoinisitol, galactitol, glycerol and the like, including cyclitols such as inositol; polyethylene glycol; amino acid polymers; sulfur-containing reducing agents, such as urea, glutathione, thioctic acid, sodium thioglycolate, thioglycerol, α -monothioglycerol and sodium thiosulfate; low molecular weight polypeptides (i.e. <10 residues); proteins such as human serum albumin, bovine serum albumin, gelatin or immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; monosaccharides such as xylose, mannose, fructose and glucose; disaccharides such as lactose, maltose and sucrose; trisaccharides such as raffinose, and polysaccharides such as dextran. Stabilizers are typically present in the range of from 0.1 to 10,000 parts by weight based on the active protein weight.

Non-ionic surfactants or detergents (also known as "wetting agents") may be present to help solubilize the therapeutic agent as well as to protect the therapeutic polypeptide against agitation-induced aggregation, which also permits the formulation to be exposed to shear surface stress without causing denaturation of the polypeptide. Suitable non-ionic surfactants include polysorbates (20, 80, etc.), polyoxamers (184, 188 etc.), Pluronic® polyols, polyoxyethylene sorbitan monoethers (Tween®-20, Tween®-80, etc.).

Additional miscellaneous excipients include bulking agents or fillers (e.g. starch), chelating agents (e.g. EDTA), antioxidants (e.g., ascorbic acid, methionine, vitamin E) and cosolvents.

The active ingredient may also be entrapped in microcapsules prepared, for example, by coascervation techniques or by interfacial polymerization, for example hydroxymethylcellulose, gelatin or poly-(methylmethacrylate) microcapsules, in colloidal drug delivery systems (for example liposomes, albumin microspheres, microemulsions, nano-particles and nanocapsules) or in macroemulsions. Such techniques are disclosed in Remington's Pharmaceutical Sciences, *supra*.

Parenteral formulations to be used for *in vivo* administration must be sterile. This is readily accomplished, for example, by filtration through sterile filtration membranes.

Sustained release preparations

Suitable examples of sustained-release preparations include semi-permeable matrices of solid hydrophobic polymers containing the polypeptide or conjugate, the matrices having a suitable form such as a film or microcapsules. Examples of sustained-release matrices include polyesters, hydrogels (for example, poly(2-hydroxyethyl-methacrylate) or poly(vinylalcohol)), polylactides, copolymers of L-glutamic acid and ethyl-L-glutamate, non-degradable ethylene-vinyl acetate, degradable lactic acid-glycolic acid copolymers such as the ProLease® technology or Lupron Depot® (injectable microspheres composed of lactic acid-glycolic acid copolymer and leuprolide acetate), and poly-D-(-)-3-hydroxybutyric acid. While polymers such as ethylene-vinyl acetate and lactic acid-glycolic acid enable release of molecules for long periods such as up to or over 100 days, certain hydrogels release proteins for shorter time periods.

When encapsulated polypeptides remain in the body for a long time, they may denature or aggregate as a result of exposure to moisture at 37°C, resulting in a loss of biological activity and possible changes in immunogenicity. Rational strategies can be devised for stabilization depending on the mechanism involved. For example, if the aggregation mechanism is discovered

to be intermolecular S-S bond formation through thio-disulfide interchange, stabilization may be achieved by modifying sulfhydryl residues, lyophilizing from acidic solutions, controlling moisture content, using appropriate additives, and developing specific polymer matrix compositions.

5 BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows purified wild-type human APC as well as various conjugates and variants of the invention. The proteins migrates on the gel as three dominate bands corresponding to the α - and β -bands of the heavy chain, with an apparent molecular weight of 41,000 and 37,000 respectively, and the light chain with an apparent molecular weight of 22,000. The degree of
10 glycosylation can also be analysed from the gel shown in Fig. 1 as the migration of the heavy chains of the conjugates D214N and M338N shifted to more cathodal positions (contrary to the variants K251N and S252N which apparently did not utilize their introduced glycosylation site) showing that these two variants are glycosylated and the site is fully utilized. From the examination of the mobility of the heavy chain subforms (α and β), it is evident that the molecular
15 weight of the carbohydrate side chains at each site is about 3,000 to 4,000.

Figure 2 shows the residual amidolytic activity of various conjugates and variants of the invention after incubation with different concentrations of alpha-1-antitrypsin (16.6 μ M (black bars) and 42.3 μ M (white bars)) for 20 hours at 37°C. Details are given in Example 11 herein.

20 *Figures 3 and 4* show the residual amidolytic activity of various conjugates and variants of the invention as a function of time in human plasma. Details, including the calculated *in vitro* half-lives in human plasma, are given in Example 13 herein.

The invention is further illustrated by the following, non-limiting, examples.

25 METHODS

Accessible Surface Area (ASA)

The computer program Access (B. Lee and F.M.Richards, J. Mol.Biol. 55: 379-400 (1971)) version 2 (©1983 Yale University) is used to compute the accessible surface area (ASA) of the individual atoms in the structure. This method typically uses a probe-size of 1.4Å
30 and defines the Accessible Surface Area (ASA) as the area formed by the center of the probe. Prior to this calculation all water molecules and all hydrogen atoms are be removed from the coordinate set. Other atoms not directly related to the protein are also removed

Fractional ASA of side chain

The fractional ASA of the side chain atoms is computed by division of the sum of the ASA of the atoms in the side chain by a value representing the ASA of the side chain atoms of that residue type in an extended ALA-x-ALA tripeptide as described in Hubbard, Campbell & Thornton (1991) J.Mol.Biol. 220, 507-530. For this example the CA atom is regarded as a part of the side chain of glycine residues but not for the remaining residues. The following values are used as standard 100% ASA for the side chain:

Ala	69.23	Å ²	Leu	140.76	Å ²
Arg	200.35	Å ²	Lys	162.50	Å ²
Asn	106.25	Å ²	Met	156.08	Å ²
Asp	102.06	Å ²	Phe	163.90	Å ²
Cys	96.69	Å ²	Pro	119.65	Å ²
Gln	140.58	Å ²	Ser	78.16	Å ²
Glu	134.61	Å ²	Thr	101.67	Å ²
Gly	32.28	Å ²	Trp	210.89	Å ²
His	147.00	Å ²	Tyr	176.61	Å ²
Ile	137.91	Å ²	Val	114.14	Å ²

Residues not detected in the structure are defined as having 100% exposure as they are thought to reside in flexible regions.

Determining distances between atoms

The distance between atoms is determined using molecular graphics software, e.g. InsightII v. 98.0, MSI Inc.

EXAMPLES*Example 1 – Determination of surface-exposed amino acids*

The coordinates for the X-ray structure of wild-type human APC (Mather, T., Oganessian, V., Hof, P., Huber, R., Foundling, S., Esmon, C., Bode, W., 1996) are available from the Protein Data Bank (PDB) (Bernstein et.al. J. Mol. Biol. (1977) 112 pp. 535) and electronically available via The Research Collaboratory for Structural Bioinformatics PDB at <http://www.pdb.org/> under accession code 1AUT. All water molecules as well as the covalently

bound inhibitor were removed from the structure before the calculation of accessible surface area was done. In the present example the betahydroxy-ASP (AP) at position 71 is treated as a normal ASP residue. The residues K156–R169 (the Lys-Arg dipeptide and the activation peptide) were not included in the calculations.

5

Sequence numbering

The sequence numbering used in this example is identical to the sequence numbering of the zymogen protein C having the amino acid sequence SEQ ID NO:4.

10

Surface exposure

Performing fractional ASA calculations on the APC molecule resulted in the following residues having zero side chain accessibility: G67, C89, C98, G103, C105, H107, C109, Y124, G142, G173, V186, L187, A198, V199, I201, V206, L207, T208, A210, C212, V221, E235, I258, A259, L260, L261, L263, A267, V274, I276, L283, V297, C331, M335, A346, G361, M364, T371, F373, L374, G376, L377, V392, I403.

15

The following residues were found to have more than 25% of their side chain exposed to the surface: Q49, L51, V52, P54, L55, E56, H57, P58, C59, A60, S61, G65, H66, T68, I70, D71, G72, I73, G74, S75, F76, S77, D79, R81, S82, G83, W84, E85, R87, F88, Q90, R91, E92, F95, L96, N97, S99, L100, D101, L110, E111, E112, V113, G114, W115, R117, S119, P122, G123, K125, G127, D128, D129, L130, L131, Q132, H134, P135, A136, V137, K138, R143, W145, K146, D172, K174, M175, R177, R178, D180, D189, S190, K191, K192, K193, H202, P203, H211, D214, E215, S216, K217, K218, L220, R229, R230, W231, K233, W234, L236, D237, D239, K241, E242, V243, F244, V245, P247, N248, S250, K251, S252, T253, T254, D255, A264, Q265, P266, T268, S270, Q271, D280, S281, G282, E285, R286, E287, Q290, A291, G292, Q293, E294, L296, Y302, H303, S304, S305, R306, E307, K308, E309, A310, K311, R312, N313, R314, T315, F316, F320, K322, P327, H328, N329, E330, S332, E333, V334, S336, N337, M338, S340, E341, I348, L349, G350, D351, R352, E357, S367, H369, G370, E382, G383, C384, L386, L387, H388, R398, D401, H404, G405, H406, R408, D409. As it appears, the active site histidine (H211) was found to be surface exposed. H211 is, however, not a candidate for being modified according to the present invention. Furthermore, the cysteine residues listed above are normally not candidates for being modified according to the present invention.

30

The following residues were found to have more than 50% of their side chain exposed to the surface: Q49, L51, V52, P54, L55, E56, A60, S61, G65, I70, D71, G72, I73, G74, S75, S77, D79, R81, S82, R87, R91, E92, F95, L96, N97, S99, E111, V113, G114, W115, R117, P122, K125, D128, D129, L130, Q132, H134, V137, K138, K146, D172, K174, R177, S190, K191, K192, K193, D214, E215, K217, K218, R229, R230, W231, K233, D239, K241, E242, P247, N248, K251, S252, Q265, P266, T268, Q271, S281, E285, Q290, G292, Y302, S305, R306, E307, K308, E309, A310, R312, N313, T315, K322, N329, E330, E333, S336, N337, M338, E341, I348, G350, R352, E357, G370, G383, H388, R398, D401, H404, G405, R408, D409.

The residues A1, N2, S3, F4, L5, E6, E7, L8, R9, H10, S11, S12, L13, E14, R15, E16, C17, I18, E19, E20, I21, C22, D23, F24, E25, E26, A27, K28, E29, I30, F31, Q32, N33, V34, D35, D36, T37, L38, A39, F40, W41, S42, K43, H44, V45, D46, G47, D48, R147, M148, E149, K150, K151, R152, S153, H154, L155, K410, E411, A412, P413, Q414, K415, S416, W417, A418, P419 are not included in the structure and are, in the present application, regarded as being 100% exposed to the surface.

Example 2 – Determination of active site region

In determining the active site region the following approach was followed: By superimposing the heavy chain of APC (1AUT) onto the X-ray structure of a ternary complex between Factor VIIa, Tissue Factor and a variant of BPTI bound in the active site (PDB accession code 1FAK. See Zhang, E., St Charles, R., Tulinsky, A.: Structure of Extracellular Tissue Factor Complexed with Factor VIIa Inhibited with a Bpti Mutant *J.Mol.Biol.* 285 pp. 2089 (1999)) using the program Modeller '98 enabled the definition of the "active site region" as any residue in the APC heavy chain having an atom within a distance of 12Å from the superimposed BPTI molecule. Furthermore, from a visual inspection a loop just outside this region (residues 306-314) was also considered to constitute part of the active site region.

Using this approach the following amino acid residues were found to be included in the "active site region":

L170, I171, D172, G173, Q184, V185, V186, L187, L188, D189, S190, K191, K192, K193, L194, A195, C196, G197, A198, T208, A209, A210, H211, C212, M213, D214, E215, S216, K217, K218, L219, L220, L228, I240, V243, V245, N248, Y249, S250, K251, S252, T253, T254, D255, N256, D257, I258, A259, L261, T295, L296, V297, T298, G299, W300, G301, Y302, H303, S304, S305, R306, E307, K308, E309, A310, K311, R312, N313, R314,

T315, F316, I321, I323, P324, V326, C331, V334, M335, S336, N337, M338, V339, M343, L344, C345, A346, G347, I348, L349, D351, R352, Q353, D354, A355, C356, E357, G358, D359, S360, G361, G362, P363, M364, G376, L377, V378, S379, W380, G381, E382, G383, C384, G385, L386, L387, H388, N389, Y390, G391, V392, Y393 and T394. Although listed
 5 here, the active site residues (H211, D257 and S360) are not candidates for being modified according to the present invention. Furthermore, the cysteine residues listed above are normally not candidates for being modified according to the present invention.

Example 3 – Determination of surface-exposed amino acids within the active site region

10 Combining the list of amino acids having more than 25% of their side chain exposed to the surface (from Example 1) with the list of amino acids included in the active site region (from Example 2), the following amino acid residues were found to be within the active site region and, at the same time, having at least 25% of its side chain exposed to the surface:

D172, D189, S190, K191, K192, K193, D214, E215, S216, K217, K218, H211, L220,
 15 V243, V245, N248, S250, K251, S252, T253, T254, D255, L296, Y302, H303, S304, S305, R306, E307, K308, E309, A310, K311, R312, N313, R314, T315, F316, V334, S336, N337, M338, I348, L349, D351, R352, E357, E382, G383, C384, L386, L387 and H388. Although listed here, the active site histidine (H211) is not a candidate for being modified according to the present invention. Moreover, C384 is normally not a candidate for being modified accord-
 20 ing to the present invention.

Example 4 - Construction of protein C expression vector

A gene encoding the human protein C precursor was constructed by assembly of synthetic oligonucleotides by PCR using methods similar to the ones described in Stemmer et al.
 25 (1995) *Gene* 164, pp. 49-53. The native protein C signal sequence was maintained in order to allow secretion of the gene product. The synthetic gene was designed with a NheI site at the 5'-end and a XbaI site at the 3'-end and subcloned behind the CMV promoter in pcDNA3.1/Hygro (Invitrogen) using these sites. The protein C precursor sequence in the resulting plasmid, termed pCR4, is given in SEQ ID NO:1.

30 Furthermore, in order to test for a higher gene expression, the synthetic gene was cloned into the KpnI-XbaI sites of pcDNA3.1/Hygro containing an intron (from pCI-Neo (Promega)) in the 5' untranslated region of the gene. The resulting plasmid was termed pRC2.

Example 5 – Site directed mutagenesis

All the mutants of protein C were constructed using Quick-Change (Stratagene). Primers were purchased from TAG Technology (Copenhagen) containing the appropriate mutations. The PCR reactions were performed according to the manufacturer's manual and the plasmids
5 were transformed into TG1 competent cells. Plasmid preparations were made on single clones and the sequences were verified using a DNA sequencer 3100 genetic Analyser (ABI)

*Primers:***D172N**

10 POF003:

CAAGTAGATCCGCGGCTCATTAACGGGAAGATGACCAGGCGGGG

POF004:

CCCCGCCTGGTCATCTTCCCGTTAATGAGCCGCGGATCTACTTG

D214N

15 EKO001:

CTGACAGCGGCCCACTGCATGAACGAGTCCAAGAAGCTCCTTGTC

EKO002:

GACAAGGAGCTTCTTGGACTCGTTCATGCAGTGGGCCGCTGTCAG

D214A

20 EKO048:

CTGACAGCGGCCCACTGCATGGCCGAGTCCAAGAAGCTCCTTGTC

EKO049:

GACAAGGAGCTTCTTGGACTCGGCCATGCAGTGGGCCGCTGTCAG

K251N

25 EKO003:

CTTCGTCCACCCCAACTACAGCAACAGCACCAACCGACAATGACATC

EKO004:

GATGTCATTGTCGGTGGTGCTGTTGCTGTAGTTGGGGTGGACGAAG

S252N

30 EKO005:

CGTCCACCCCAACTACAGCAAGAACACCAACCGACAATGACATCGC

EKO006:

GCGATGTCATTGTCGGTGGTGTTCTTGCTGTAGTTGGGGTGGACG

Y302N

EKO007:

CCCTCGTGACGGGCTGGGGCAACCACAGCAGCCGAGAGAAGGAGGCC

EKO008:

5 GGCCTCCTTCTCTCGGCTGCTGTGGTTGCCCCAGCCCGTCACGAGGG

M338N

EKO011:

CAGCGAGGTCATGAGCAACAACGTGTCTGAGAACATGC

EKO012:

10 GCATGTTCTCAGACACGTTGTTGCTCATGACCTCGCTG

M338A

EKO046:

GCAGCGAGGTCATGAGCAACGCCGTGTCTGAGAACATGC

EKO047:

15 GCATGTTCTCAGACACGGCGTTGCTCATGACCTCGCTGC

D189N+K191N

EKO019:

CCCCTGGCAGGTGGTCCTGCTGAACTCAAACAAGAAGCTGGCCTGCGGGG

EKO020:

20 CCCC GCAGGCCAGCTTCTTGTTTGAGTTCAGCAGGACCACCTGCCAGGGG

D189N+K191T

EKO033:

CCCCTGGCAGGTGGTCCTGCTGAACTCAACCAAGAAGCTGGCCTGCGGGG

EKO034:

25 CCCC GCAGGCCAGCTTCTTGTTTGAGTTCAGCAGGACCACCTGCC

S190N+K192T

EKO044: GGCAGGTGGTCCTGCTGGACAACAAGACCAAGCTGGCCTGCGGGGCAG-TGC

EKO045: GCACTGCCCCGCAGGCCAGCTTGGTCTTGTTGTCCAGCAGGACCACCT-

30 GCC

K191N+K193T

EKO050:

GTCCTGCTGGACTCAAACAAGACCCTGGCCTGCGGGGCAGTG

EKO051:

CACTGCCCCGCAGGCCAGGGTCTTGTTTGAGTCCAGCAGGAC

K217N+L219T

EKO029:

5 GCATGGATGAGTCCAACAAGACCCTTGTCAGGCTTGGAGAGTATGACC

EKO030:

GGTCATACTCTCCAAGCCTGACAAGGGTCTTGTTGGACTCATCCATGC

T253N+D255T

EKO031: CCAACTACAGCAAGAGCAACACCACCAATGACATCGCACTGCTGCACCT-

10 GGC

EKO032: GCCAGGTGCAGCAGTGCGATGTCATTGGTGGTGTTGCTCTTGCTGTAG-
TTGG

S305N+E307T

EKO023:

15 GGCTGGGGCTACCACAGCAACCGAACCAAGGAGGCCAAGAGAAACCGC

EKO024:

GCGGTTTCTCTTGGCCTCCTTGGTTCGGTTGCTGTGGTAGCCCCAGCC

E307N+E309T

EKO025:

20 GGCTACCACAGCAGCCGAAACAAGACCGCCAAGAGAAACCGCACCTTCG

EKO026:

CGAAGGTGCGGTTTCTCTTGGCGGTCTTGTTTCGGCTGCTGTGGTAGCC

S336N+M338T

EKO027:

25 GCAGCGAGGTCATGAACAACACCGTGTCTGAGAACATGCTGTGTGCGGG

EKO028:

CCCGCACACAGCATGTTCTCAGACACGGTGTTGTTTCATGACCTCGCTGC

L386N+H388T

EKO017: GGTGAGCTGGGGTGAGGGCTGTGGGAACCTTACCAACTACGGCGTTTA-

30 CACC

EKO018: GGTGTAAACGCCGTAGTTGGTAAGGTTCCCACAGCCCTCACCCCAGCT-
CACC

Example 6 – Production

Transient expression of wild-type protein C and protein C variants was performed using the Fugene transfection reagent (Roche) in COS 7 cells grown in DMEM (Gibco 21969-035) supplemented with 10% fetal serum, 2 mM L-glutamine, 100 U/ml of penicillin, 100
5 μ g/ml streptomycin and 5 μ g/ml vitamin K. On the day of transfection the medium was substituted with fresh medium 4-5 hours before transfection. The day after transfection the medium was substituted with serum-free production medium based on DMEM (Gibco 31053-028) supplemented with 2 mM L-glutamine, 1 mM Sodium Pyruvate, 1/500 Ex-cyte (serologicals) 1/100 ITSA (Gibco 51300-044), 100 U/ml of penicillin, 100 μ g/ml streptomycin and 5 μ g/ml
10 vitamin K. After incubation for two days the medium was harvested and the expressed variants were analysed for production and activity (see Example 9 below).

Example 7- Purification

Approximately 15 mg Ca specific monoclonal antibody was coupled to 5 ml CNBr-
15 activated Sepharose FF from Pharmacia according to the manufacturer's instructions. Approximately 1 ml of the coupled matrix was packed in a HR 10 column and washed with buffer A (20 mM Tris, 0.3 M NaCl, 5 mM CaCl₂, pH 7.5) at a flow rate of 1 ml/min. Approximately 90 ml of sterile filtered culture medium was made 0.3 M NaCl and 5 mM CaCl₂ and applied to the column at the same flow rate. Prior to elution, the column was washed with 20 column vol-
20 umes of buffer A. Elution was carried out with buffer B (20 mM Tris and 10 mM EDTA, pH 7.5) and fractionated in 1 ml fractions. Fractions containing protein C, as judged by OD₂₈₀, western blot and SDS-PAGE, were pooled and stored at -80°C. The above-described purification procedure represents one out of several possible procedures for purifying protein C (see, for example, Kiesel, J. Clin. Invest. (1979) 64 pp. 761-769).

25 The purified proteins were activated using the activation protocol (see Example 8 below). The purity of all proteins was checked using polyacrylamide gel electrophoresis (PAGE) analysis. In addition, the degree of glycosylation was estimated from these gel analyses by monitoring changes in molecular weight. Increased apparent molecular weights compared to the wild-type human APC molecule demonstrate that the APC variants have been glycosylated.

30 An example of the wild-type APC and APC variants can be seen in Figure 1. The proteins migrate on the gel as three dominant bands corresponding to the α - and β -bands of the heavy chain, with an apparent molecular weight of 41,000 and 37,000 respectively, and the light chain with an apparent molecular weight of 22,000. The degree of glycosylation was also

investigated in the PAGE analysis. Figure 1 includes two APC variants that are glycosylated in the introduced glycosylation site. The migration of the heavy chains of the APC variant D214N and M338N shifted to more cathodal positions, showing that these two variants are glycosylated and the site is fully used. From the examination of the mobility of the heavy chain subforms (α and β), it is evident that the molecular weight of the carbohydrate side chains at each site is about 3,000 to 4,000.

Example 8 - Activation

The protein C variants and conjugates were activated using the venom protein C activator, ACC-C (Nakagaki et al., Thrombosis Research 58:593-602, 1990). The zymogen forms were incubated at 37°C for about 60 min in 50 mM Tris-HCl (pH 7.5), 100 mM NaCl, 5 mM EDTA, using a final concentration of 1 ng/ml of ACC-C. The activation process was checked using the APC amidolytic activity assay (see example 9 below) and polyacrylamide gel electrophoresis analysis.

Example 9 – Determination of amidolytic activity

APC Amidolytic Assay

The amidolytic activity of human APC and the compounds of the invention is determined using the peptide substrate SPECTROZYME PCa with the formula H-D-Lys(γ -Cbo)-Pro-Arg-pNA.2AcOH (American Diagnostica Inc, product # 336) at a final concentration of 0.5 mM. Assays are performed at 23°C in 50 mM Tris-HCl (pH 8.3), 100 mM NaCl, 5 mM CaCl₂. The rate of hydrolysis of the PCa substrate by human APC and the compounds of the invention are recorded for 3 min at 405 nm as the change in absorbance units/min in a plate reader.

Results

All expressed and activated conjugates and variants were analysed for activity. 4 μ l (unpurified) cell culture medium was assayed as described right above. The obtained activities, which do not reflect the specific activities since they depend *inter alia* on the expression level, indicate whether the proteins were expressed and whether they possessed activity.

The following activities were obtained:

Table 1a

<u>Compound</u>	<u>milliOD₄₀₅/min</u>
-----------------	----------------------------------

		59
	wild-type COS 7 APC	41
	D214N	28
	D214A (control)	10
	K251N*	19
5	S252N*	16
	Y302N*	14
	M338N	53
	M338A (control)	33
	D189N+K191T	8
10	D189N+K191N (control)	12
	S190N+K192T*	29
	K191N+K193T	4
	K217+L219T	16
	T253N+D255T	2
15	S305N+E307T	6
	E307N+E309T	30
	S336N+M338T	4
	L386N+H388T	13

*: No detectable sugar moiety attached to the introduced glycosylation site as judged from SDS-PAGE

20

Selected candidates were purified and their specific amidolytic activities were measured in the above assay using a protein concentration of 30 nM. The following activities were found:

25 *Table 1b*

	Compound	milliOD ₄₀₅ /min	% of wild-type APC
	wild-type COS 7 APC	48.9	-
	D214N	34.8	71
	K251N*	45.2	92
30	S252N*	43.1	88
	M338N	44.8	92
	S336N+M338T	41.5	85
	L386N+H388T	23.0	47

*: No detectable sugar moiety attached to the introduced glycosylation site as judged from SDS-PAGE

As it appears, the specific amidolytic activity of the tested conjugates and variants is at the same level as the wild-type human APC molecule.

5 Example 10 – Determination of anticoagulant activity

APC Clotting Assay

Anticoagulant activity is assessed by monitoring the prolongation of clotting time in the activated partial thromboplastin time (APTT) assay using Nycoplastin (Nycomed, product no. 1002448) together with Normal Hemostasis Reference Plasma (American Diagnostica Inc.,
10 catalogue no. 258N). Coagulation is started by mixing the APTT reagent containing human APC or compounds of the invention with the normal hemostasis reference plasma at 37°C and measuring the clotting time by manual mixing. The clotting time for the human APC is compared to the clotting time of the compounds of the invention to calculate the anticoagulant activity expressed in percentage to the human APC anticoagulant activity.

15

Results

Using the above assay the following anticoagulant activities were found:

Table 2

20	<u>Compound</u>	<u>Anticoagulant activity (% of human APC)</u>
	D214N	22.4
	K251N*	24.5
	S252N*	24.5
	M338N	34.7
25	<u>L386N+H388T</u>	<u>14.3</u>

*: No detectable sugar moiety attached to the introduced glycosylation site as judged from SDS-PAGE

These results show that the anticoagulant properties of the conjugates and variants of the invention are preserved to a large extent. This clearly shows that it is possible to design
30 APC variants and conjugates with significantly increased resistance toward inhibition in plasma (see examples below) with retained anticoagulant activity.

Example 11 – Inactivation by alpha-1-antitrypsin

Alpha-1-Antitrypsin Inactivation Assay

Human APC or compounds of the invention are incubated with 16.6 or 42.3 μ M human alpha-1-antitrypsin (Sigma) in 10 mM Tris-HCl (pH 7.5), 150 mM NaCl, 5 mM CaCl₂ containing 0.1% BSA at 37°C. After 20 hours incubation a 15 μ l sample of the incubated mix-
 5 tures is added to 110 μ l 50 mM Tris-HCl (pH 8.3), 100 mM NaCl, 5 mM CaCl₂ in microplates and assayed for APC amidolytic activity as described in the "APC Amidolytic Assay". The remaining activity is calculated by normalizing with the activity obtained in samples lacking alpha-1-antitrypsin but otherwise incubated under identical conditions.

10 *Results*

Using the above assay the following results were obtained:

Table 3

	Compound	% residual amidolytic activity	
		16.6 μ M inhibitor	42.3 μ M inhibitor
15	wild-type plasma APC	10	2
	wild-type COS 7 APC	7	<1
	D214N	80	81
	D214A (control)	21	1
20	K251N*	62	53
	S252N*	62	34
	Y302N*	50	30
	M338N	38	12
	M338A (control)	9	2
25	D189N+K191T	90	77
	D189N+K191N (control)	12	<1
	S190N+K192T*	28	5
	K191N+K193T	59	24
	K217+L219T	20	4
30	T253N+D255T	56	38
	S305N+E307T	42	9
	E307N+E309T	10	<1
	S336N+M338T	72	40

L386N+H388T

68

44

*: No detectable sugar moiety attached to the introduced glycosylation site as judged from SDS-PAGE

The data are also shown in Fig.2. The results show that practically all of the conjugates have increased resistance towards alpha-1-antitrypsin inhibition. In particular, D214N and D189N+K191T retain more than 70% of their amidolytic activity even at the highest alpha-1-antitrypsin concentration. The effect of the glycosylation of these compounds can be seen when comparing these two conjugates with D214A and D189N+K191N, which lack glycosylation. These variants are inhibited significantly more than their glycosylated equivalents indicating that glycosylation is important for improving the resistance towards alpha-1-antitrypsin inhibition. Moreover, it should be noted that the variants K251N, S252N, Y302N and S190+K192T, which apparently have not utilized their introduced glycosylation site (as judged from SDS-PAGE), have significantly increased their resistance towards alpha-1-antitrypsin inhibition as compared to wild-type human APC.

Example 12 – Inactivation by human plasma

Human Plasma Inactivation Assay I

Human APC or compounds of the invention are incubated in 90% normal human plasma (Sigma Diagnostics, AccucutTM reference plasma) containing 50 mM Tris-HCl (pH 7.5), 100 mM NaCl, 5 mM CaCl₂ at 37°C. Aliquots are removed after 200 min and assayed for APC amidolytic activity as described in the “APC Amidolytic Assay”. The residual APC activity after 200 min is expressed in percentage of the APC activity measured at the start of the experiment.

Results

Using the above assay the following results were obtained:

Table 4

Compound	% residual amidolytic activity after 200 min in 90% normal human plasma
wild-type plasma APC	5
wild-type COS 7 APC	7
D214N	80

	K251N*	57
	S252N*	45
	M338N	22
	S336N+M338T	45
5	<u>L336N+H388T</u>	<u>72</u>

*: No detectable sugar moiety attached to the introduced glycosylation site as judged from SDS-PAGE

The above results clearly indicated that the conjugates as well as the variants according to the invention are highly resistance towards inactivation in human plasma.

10

Example 13 – In vitro half-life in human plasma

Human Plasma Inactivation Assay II

Human APC or compounds of the invention are incubated in 90% normal human plasma (Sigma Diagnostics, Accuclot™ reference plasma) containing 50 mM Tris-HCl (pH 7.5), 100 mM NaCl, 5 mM CaCl₂ at 37°C. Aliquots are removed at various time-points and assayed for APC amidolytic activity as described in the “APC Amidolytic Assay”. The residual APC activity at the various time-points is expressed in percentage of the APC activity measured at the start of the experiment. The *in vitro* half-life (expressed in minutes) is calculated as the time at which 50% of the APC activity is still present.

20

Results

The following *in vitro* half-lives were obtained:

Table 5

25	Compound	<i>In vitro</i> half-life (min)	Fold increase relative to wild-type human APC
	wild-type plasma APC	40	-
	wild-type COS 7 APC	42	-
	D214N	>400	>10
30	K251N*	255	6.4
	S252N*	155	3.9
	M338N	85	2.1
	S336N+M338T	185	4.6
	<u>L386N+H388T</u>	<u>>400</u>	<u>>10</u>

*: No detectable sugar moiety attached to the introduced glycosylation site as judged from SDS-PAGE

The experimental data points are shown in Figs. 3 and 4. The results show that the APC variants and conjugates have significantly increased *in vitro* half-lives in human plasma.

- 5 Especially the D214N and L386N+H388T conjugates show a significantly increased *in vitro* half-life (increased more than 10 times).

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WO 02/32461

PCT/DK01/00679

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WO 02/32461

PCT/DK01/00679

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WO 02/32461

PCT/DK01/00679

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WO 02/32461

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WO 02/32461

PCT/DK01/00679

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WO 02/32461

PCT/DK01/00679

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Gly His Gly Thr Cys Ile Asp Gly Ile Gly Ser Phe Ser Cys Asp Cys	65	70	75
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Asp Ile Ala Leu Leu His Leu Ala Gln Pro Ala Thr Leu Ser Gln Thr	260	265	270
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WO 02/32461

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CLAIMS

1. A conjugate comprising at least one non-polypeptide moiety covalently attached to a protein C polypeptide that comprises an amino acid sequence which differs from that of a parent protein C polypeptide in at least one introduced and/or at least one removed amino acid residue comprising an attachment group for said non-polypeptide moiety.
2. The conjugate according to claim 1, wherein the parent protein C polypeptide has the amino acid sequence shown in SEQ ID NO:4 or is a variant thereof.
3. The conjugate according to claim 2, wherein the parent protein C polypeptide has the amino acid sequence shown in SEQ ID NO:4.
4. The conjugate according to any of claims 1-3 in its activated form.
5. The conjugate according to any of claims 1-4, wherein at least one attachment group for the non-polypeptide moiety has been introduced.
6. The conjugate according to claim 5, wherein at least one glycosylation site has been introduced.
7. The conjugate according to claim 6, wherein the glycosylation site is an *in vivo* N-glycosylation site.
8. The conjugate according to claim 7, wherein the glycosylation site has been introduced in a position which is occupied by an amino acid residue having at least 25% of its side chain exposed to the surface (as defined in Example 1 herein).
9. The conjugate according to claim 8, wherein the introduced glycosylation site is selected from the group consisting of D172N+K174S, D172N+K174T, D189N+K191S, D189N+K191T, S190N+K192S, S190N+K192T, K191N+K193S, K191N+K193T, K192N+L194S, K192N+L194T, K193N+A195S, K193N+A195T, D214N, D214N+S216T, E215N+K217S, E215N+K217T, S216N+K218S, S216N+K218T, K217N+L219S,

K217N+L219T, K218N+L220S, K218N+L220T, L220N+R222S, L220N+R222T,
V243N+V245S, V243N+V245T, V245N+P247S, V245N+P247T, S250N, S250N+S252T,
K251N, K251N+T253S, S252N, S252N+T254S, T253N+D255S, T253N+D255T,
T254N+N256S, T254N+N256T, D255N+D257S, D255N+D257T, L296N, L296N+T298S,
5 Y302N, Y302N+S304T, H303N, H303N+S305T, S304N+R306S, S304N+R306T,
S305N+E307S, S305N+E307T, R306N+K308S, R306N+K308T, E307N+E309S,
E307N+E309T, K308N+A310S, K308N+A310T, E309N+K311S, E309N+K311T,
A310N+R312S, A310N+R312T, R312N+R314S, R312N+R314T, T315N+V317S,
T315N+V317T, F316N+L318S, F316N+L318T, V334N, V334N+S336T, S336N+M338S,
10 S336N+M338T, V339S, V339T, M338N, M338N+S340T, I348N+G350S, I348N+G350T,
L349N+D351S, L349N+D351T, D351N+Q353S, D351N+Q353T, R352N+D354S,
R352N+D354T, E357N+D359S, E357N+D359T, G383N+G385S, G383N+G385T,
L386N+H388S, L386N+H388T, L387N+N389S, L387N+N389T, H388N+Y390S and
H388N+Y390T.

15

10. The conjugate according to claim 9, wherein the introduced glycosylation site is selected
from the group consisting of D189N+K191S, D189N+K191T, S190N+K192S, S190N+K192T,
K191N+K193S, K191N+K193T, D214N, D214N+S216T, K217N+L219S, K217N+L219T,
K251N, K251N+T253S, S252N, S252N+T254S, T253N+D255S, T253N+D255T, Y302N,
20 Y302N+S304T, T253N+D255S, T253N+D255T, S336N+M338S, S336N+M338T, V339S,
V339T, M338N, M338N+S340T, G383N+G385S, G383N+G385T, L386N+H388S and
L386N+H388T.

11. The conjugate according to claim 10, wherein the introduced glycosylation site is selected
25 from the group consisting of D189N+K191S, D189N+K191T, K191N+K193T, D214N,
D214N+S216T, K251N, K251N+T253S, S252N, S252N+T254S, T253N+D255S,
T253N+D255T, Y302N, Y302N+S304T, S305N+E307T, S305N+E307S, S336N+M338S,
S336N+M338T, V339S, V339T, M338N, M338N+S340T, G383N+G385S, G383N+G385T,
L386N+H388S and L386N+H388T.

30

12. The conjugate according to claim 11, wherein the introduced glycosylation site is selected
from the group consisting of D189N+K191T, K191N+K193T, D214N, K251N, S252N,

T253N+D255T, Y302N, S305N+E307T, S336N+M338T, V339T, M338N, G383N+G385T and L386N+H388T.

13. The conjugate according to claim 12, wherein the introduced glycosylation site is selected
5 from the group consisting of D189N+K191T, K191N+K193T, D214N, T253N+D255T, S305N+E307T, S336N+M338T, M338N, G383N+G385T and L386N+H388T.

14. The conjugate according to claim 13, wherein the introduced glycosylation site is selected from the group consisting of D189N+K191T, D214N and L386N+H388T.

10

15. The conjugate according to any of claims 1-4 wherein the introduced and/or removed attachment group is selected from the group consisting of a lysine residue, a glutamic acid residue, an aspartic acid residue, a tyrosine residue, a serine residue and a cysteine residue.

15 16. The conjugate according to claim 15, wherein the attachment group has been introduced in or removed from a position which is occupied by an amino acid residue having at least 25% of its side chain exposed to the surface (as defined in Example 1 herein).

17. The conjugate according to claim 16, wherein the attachment group is introduced in or re-
20 moved from a position selected from the group consisting of D172, D189, S190, K191, K192, K193, D214, E215, S216, K217, K218, L220, V243, V245, S250, K251, S252, T253, T254, L296, Y302, H303, S304, S305, R306, E307, K308, E309, A310, R312, T315, F316, V334, S336, N337, M338, I348, L349, D351, R352, E357, G383, L386, L387 and H388.

25 18. The conjugate according to claim 17, wherein the attachment group is introduced in or removed from a position selected from the group consisting of D189, S190, K191, D214, K217, K251, S252, T253, Y302, S305, E307, S336, N337, M338, G383 and L386.

19. The conjugate according to claim 18, wherein the attachment group is introduced in a or
30 removed from a position selected from the group consisting of D189, D214, K251, S252, T253, Y302, S305, S336, N337, M338, G383 and L386.

20. The conjugate according to any of claims 15-19, wherein the introduced attachment group is a cysteine residue.

21. The conjugate according to any of claims 15-20, wherein the non-polypeptide moiety is a
5 polymer molecule.

22. The conjugate according to claim 21, wherein the non-polypeptide moiety is a linear or branched polyethylene glycol or a polyalkylene oxide.

10 23. The conjugate according to any of claims 5-14, which further comprises a non-polypeptide moiety as defined in any of claims 15-22.

24. The conjugate according to any of claims 1-23, which, in its activated form, and when tested in the "APC Amidolytic Assay" described in Example 9 herein, has an activity of at least
15 10% of the wild-type human APC activity.

25. The conjugate according to any of claims 1-24, which, in its activated form and when tested in the "APC Clotting Assay" described in Example 10, has an anticoagulant activity of at least
20 10% of the wild-type human APC anticoagulant activity.

26. The conjugate according to any of claims 1-25, which, in its activated form, has an increased resistance towards inactivation by alpha-1-antitrypsin as compared to human APC.

27. The conjugate according to claim 26, which, in its activated form has a residual activity of
25 at least 20% when tested in the "Alpha-1-Antitrypsin Inactivation Assay" described in Example 11 herein using an inhibitor concentration of 16.6 μ M.

28. The conjugate according to any of claims 1-27, which, in its activated form, has an increased resistance towards inactivation by human plasma.

30

29. The conjugate according to claim 28, which, in its activated form and when tested in the "Human Plasma Inactivation Assay I" described in Example 12 herein, has a residual activity of at least 20%.

30. The conjugate according to claim 28 or 39, where the ratio between the *in vitro* half-life of said conjugate, in its activated form, and the *in vitro* half-life of human APC is at least 1.25 when tested in the "Human Plasma Inactivation Assay II" described in Example 13 herein.

5

31. The conjugate according to any of claims 1-30, which, in its activated form, has an increased functional *in vivo* half-life or an increased serum half-life as compared to human APC.

32. The conjugate according to claim 31, wherein the ratio between the functional *in vivo* half-life or the serum half-life of said conjugate and the functional *in vivo* half-life or serum half-life of human APC is at least 1.25.

33. A variant of a parent protein C polypeptide, said variant comprising a substitution in a position selected from the group consisting of D172, D189, S190, K191, K192, K193, D214, E215, S216, K217, K218, L220, V243, V245, S250, K251, S252, T253, T254, D255, L296, Y302, H303, S304, S305, R306, E307, K308, E309, A310, R312, T315, F316, V334, S336, N337, M338, I348, L349, D351, R352, E357, E382, G383, L386, L387 and H388, with the proviso that the substitution is not selected from the group consisting of T254S, T254A, T254H, T254K, T254R, T254N, T254D, T254E, T254G, T254Q, Y302S, Y302A, Y302T, Y302H, Y302K, Y302R, Y302N, Y302D, Y302E, Y302G, Y302Q, F316S, F316A, F316T, F316H, F316K, F316R, F316N, F316D, F316E, F316G and F316Q.

25

34. The variant according to claim 33, wherein the parent protein C polypeptide has the amino acid sequence shown in SEQ ID NO:4.

35. The variant according to claim 33 or 34 in its activated form.

36. The variant according to any of claims 33-35, wherein said variant is as the polypeptide part of the conjugate defined in any of claims 9-20.

30

37. The variant according to claim 36, wherein said variant comprises a substitution selected from the group consisting of K251N, S252N and Y302N.

38. The variant according to any of claims 33-37, which has the properties defined in any of claims 24-32.

39. A nucleotide sequence encoding the variant as defined in any of claims 33-38.

5

40. An expression vector comprising a nucleotide sequence as defined in claim 39.

41. A host cell comprising a nucleotide sequence as defined in claim 39 or an expression vector as defined in claim 40.

10

42. The host cell according to claim 41, which is selected from the group consisting of COS, CHO, BHK and HEK293 cells.

43. A pharmaceutical composition comprising a conjugate as defined in any of claims 1-32 or a variant as defined in any of claims 33-38 and a pharmaceutically acceptable carrier or excipient.

15

44. A conjugate as defined in any of claims 1-32, a variant as defined in any of claims 33-38 or a pharmaceutical composition as defined in claim 43 for use as a medicament.

45. Use of a conjugate as defined in any of claims 1-32, use of a variant as defined in any of claims 33-38, or use of a pharmaceutical composition as defined in claim 43 for the manufacture of a medicament for the treatment of stroke; myocardial infarction; after venous thrombosis; disseminated intravascular coagulation (DIC); sepsis; septic shock; emboli, such as pulmonary emboli; transplantation, such as bone marrow transplantation; burns; pregnancy; major surgery/traum or adult respiratory stress syndrome (ARDS).

25

46. The use according to claim 45 for the manufacture of a medicament for the treatment of septic shock

47. A method for treating or preventing a disease selected from the group consisting of stroke; myocardial infarction; after venous thrombosis; disseminated intravascular coagulation (DIC); sepsis; septic shock; emboli, such as pulmonary emboli; transplantation, such as bone marrow transplantation; burns; pregnancy; major surgery/traum and adult respiratory stress syndrome

30

(ARDS), the method comprising administering to a patient in need thereof an effective amount of a conjugate as defined in any of claims 1-32, or of a variant as defined in any of claims 33-38, or of a pharmaceutical composition as defined in claim 43.

5 48. The method according to claim 47 for treating or preventing septic shock.

49. A method for producing a conjugate as defined in any of claims 1-32, the method comprises culturing an appropriate host cell under conditions conducive for the expression of the polypeptide part of the conjugate, and recovering the polypeptide, wherein

10 a) the polypeptide comprises at least one N- or O-glycosylation site and the host cell is an eukaryotic host cell capable of *in vivo* glycosylation, and/or

b) the polypeptide is subjected to conjugation to a non-polypeptide moiety *in vitro*.

50. A method of increasing the functional *in vivo* half-life or the serum half-life of a parent
15 protein C polypeptide, which method comprises introducing an amino acid residue constituting an attachment group for a non-polypeptide moiety into a position of a parent protein C polypeptide comprising an amino acid residue having at least 25% of its side chain exposed to the surface (as defined in Example 1 herein) which does not contain such attachment group and/or
removing an amino acid residue constituting such attachment group, and subjecting the result-
20 ing modified polypeptide to conjugation with the non-polypeptide moiety which has the amino acid residue having been introduced and/or removed as an the attachment group.

1/4

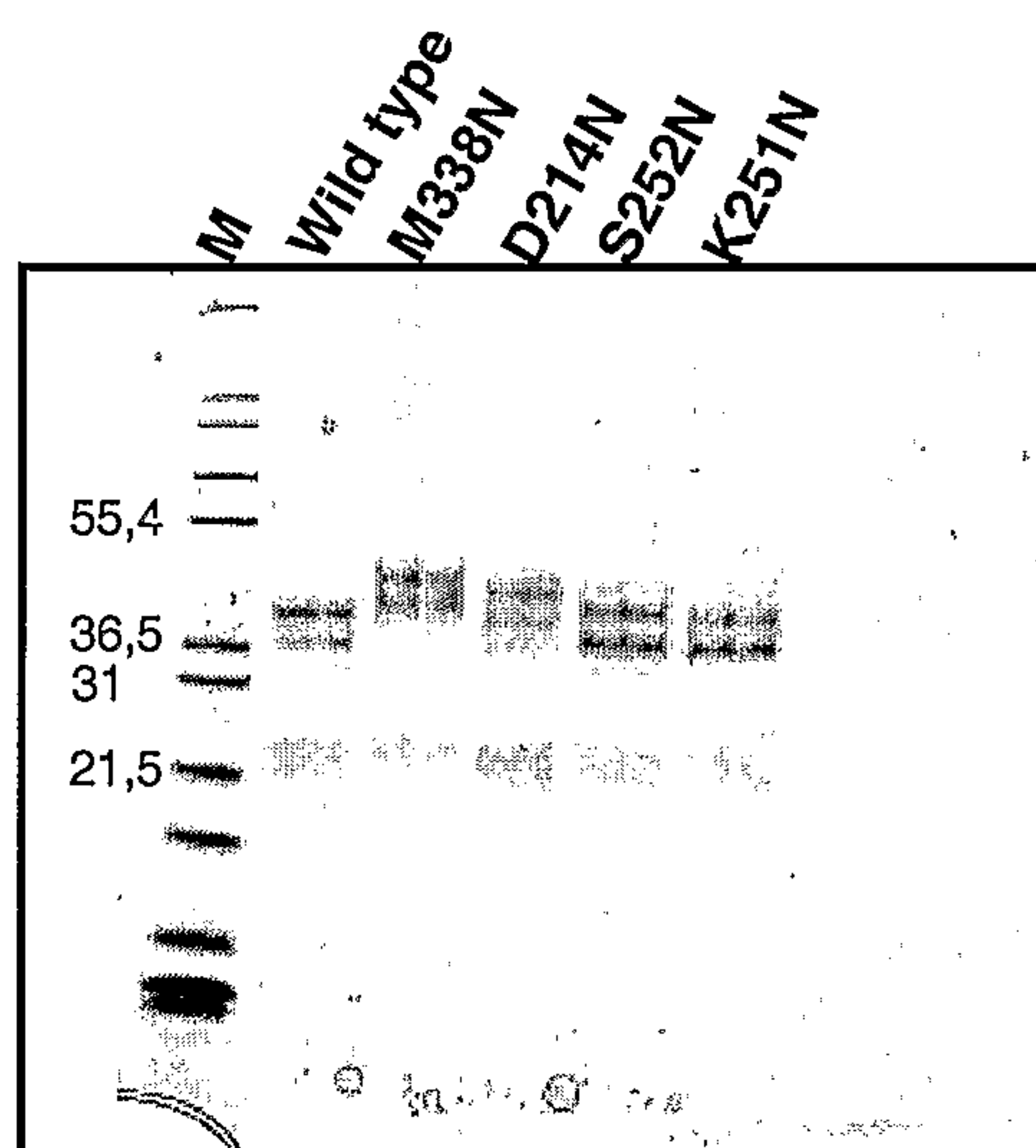


Fig. 1

2/4

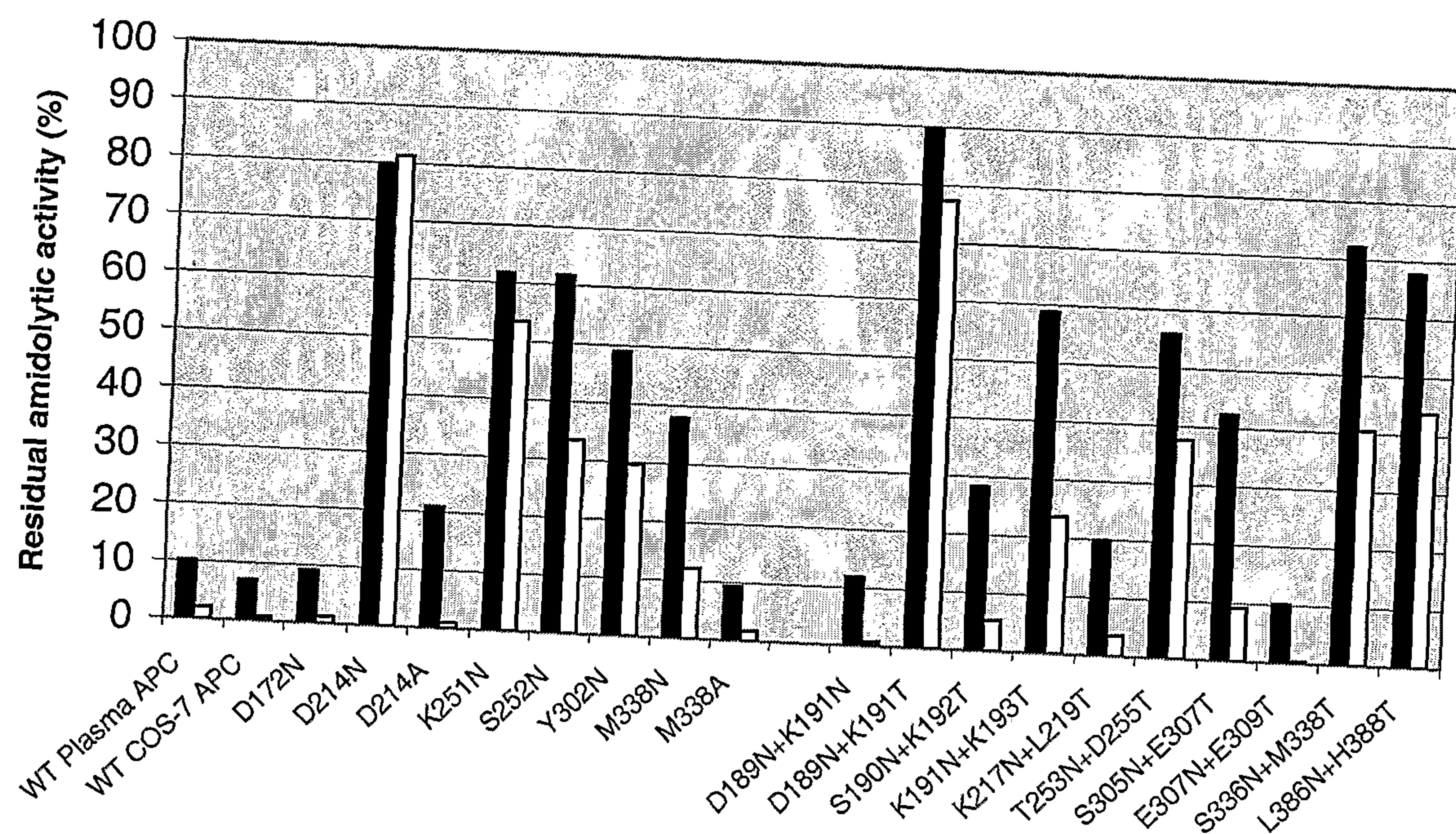


Fig. 2

3/4

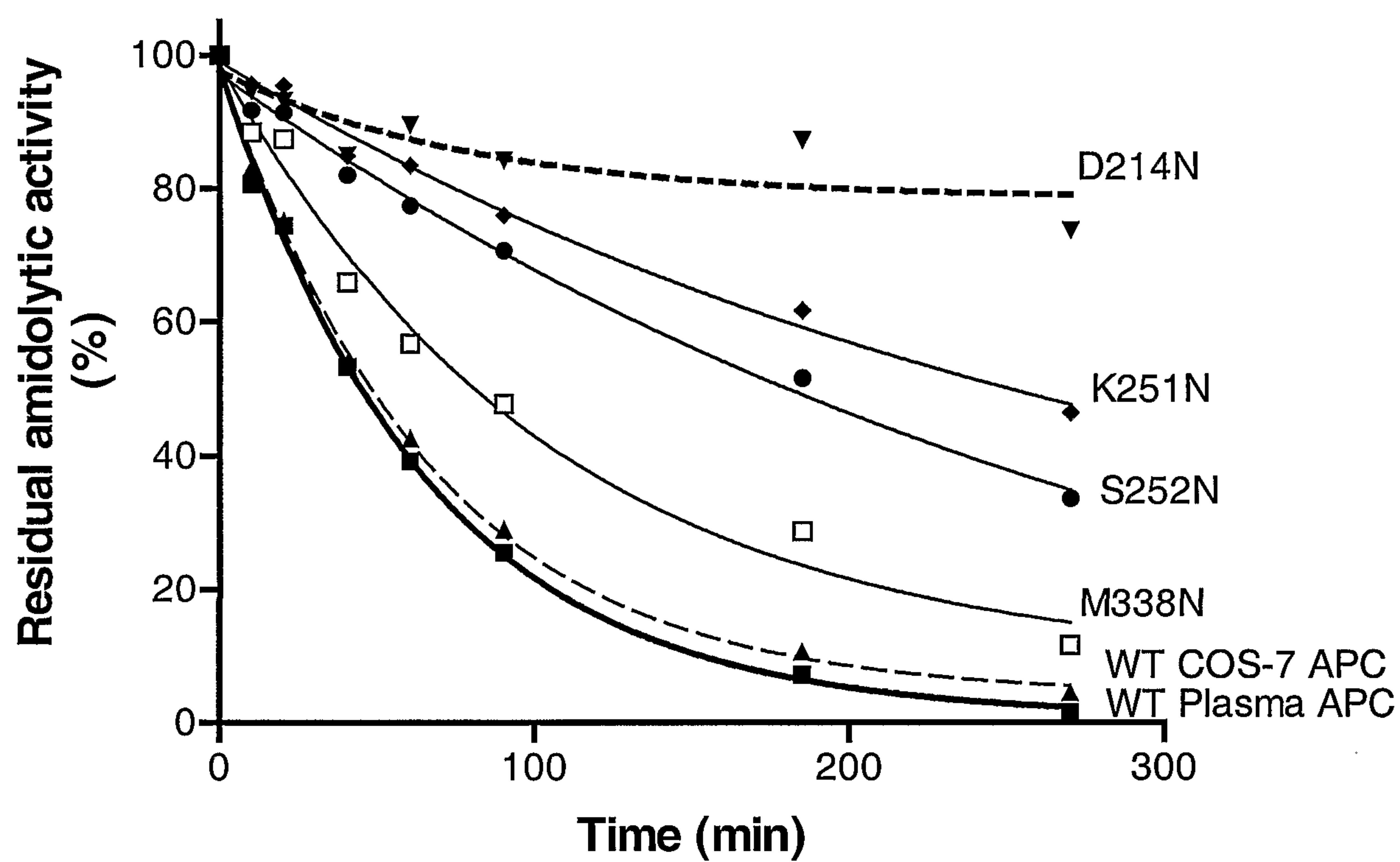


Fig. 3

4/4

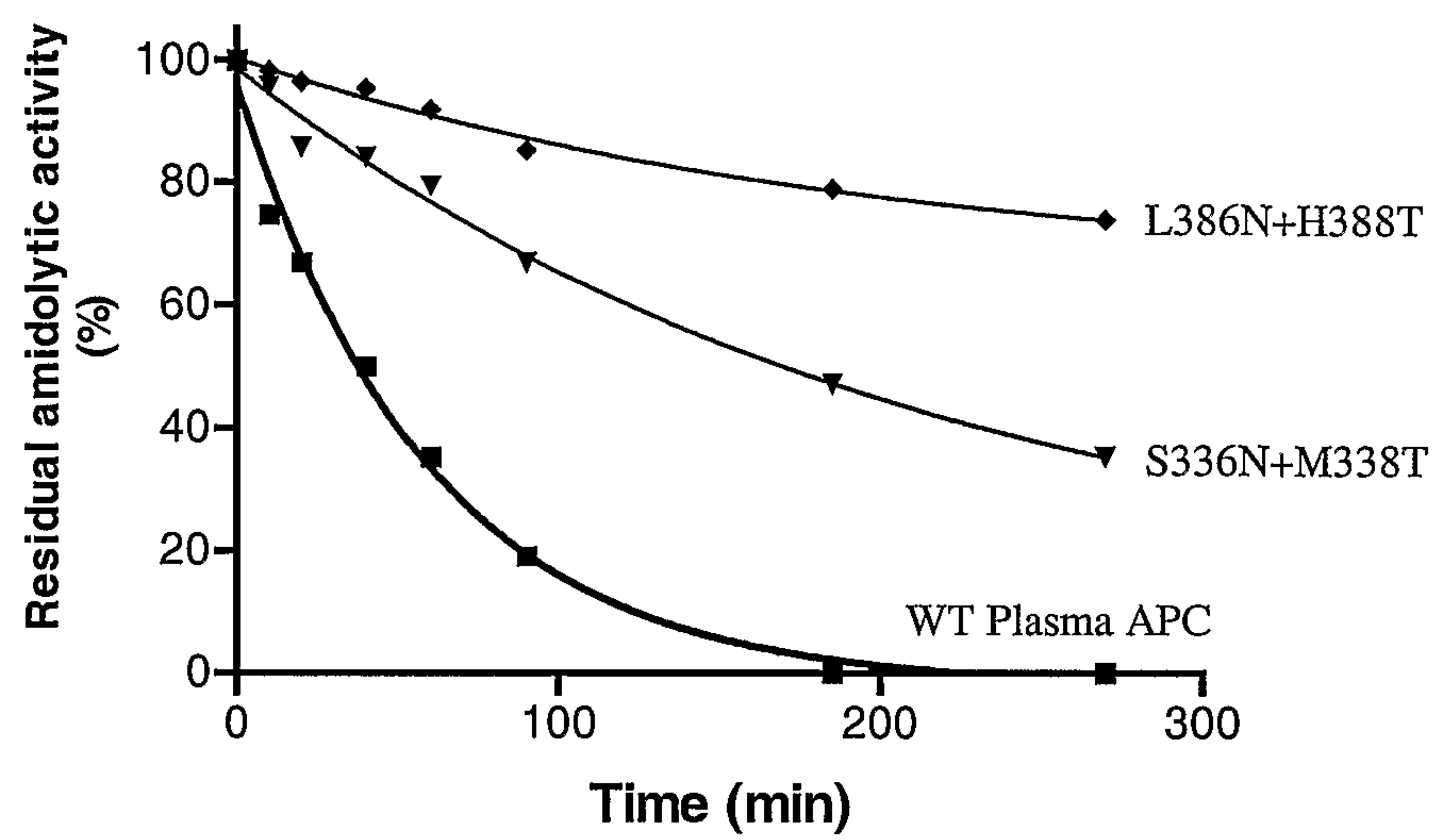


Fig. 4

